INTRODUCTION

The mediastinum is located centrally in the thorax between the lungs. It has recently been classified by the International Thymic Malignancy Interest Group (ITMIG) into the following three compartments (1): prevascular, visceral, and paravertebral. Precise and consistent compartmentalization is possible with this new model.

Mediastinal masses are rarely identified. Henschke et al. (2) reported a prevalence of 0.8% in asymptomatic people with a high risk of lung cancer. However, mediastinal masses are observed in a wide spectrum of benign and malignant diseases. As computed tomography (CT) screening for lung cancer is introduced worldwide, several mediastinal masses will be detected. Therefore, radiologists must be familiar with identifying these masses.

Nowadays, magnetic resonance imaging (MRI) is a useful diagnostic tool for evaluating mediastinal masses considering that several lesions that appear indeterminate on CT can be differentiated on MRI. In particular, MRI is a useful diagnostic tool for differentiating cystic from solid lesions, evaluating invasiveness into adjacent structures, and characterizing tissue components such as fat or hemorrhage.

When mediastinal masses are observed on CT or MRI, the first step is to localize the lesion into one of the three compartments, each of which has its own list of differential diagnoses. The next step is to analyze imaging findings in detail. With a basic knowledge of MRI, radiologists can easily diagnose mediastinal masses (3). Here, we review various MRI findings in mediastinal masses localized in each of the three compartments of the ITMIG classification system.
Basic Principles and Benefits of Magnetic Resonance Imaging

MRI has excellent contrast resolution. It is specifically useful for evaluating cystic lesions and identifying fat within mediastinal masses. The most frequently encountered type of mass on mediastinal MRI is a cystic lesion such as a thymic, bronchogenic, or pericardial cyst. Cysts show high signal intensity on T2-weighted images (4) and occasionally show high or intermediate signal intensity on T1-weighted images due to hemorrhagic or proteinaceous components (4, 5). A focus of enhancement within a cystic lesion is

Fig. 1. A 42-year-old woman with a pericardial inflammatory myofibroblastic tumor.
An axial contrast-enhanced computed tomography image at the level of the ascending thoracic aorta (A) shows a heterogeneously enhancing soft tissue mass (asterisk) in the thymic bed of the prevascular mediastinum. Initially, it was considered that the most likely diagnosis was a thymic epithelial tumor. MR imaging was performed for further evaluation; sagittal cine imaging (B) shows the intrapericardial location of the mass. Surgical excision was performed, and the histological diagnosis was an inflammatory myofibroblastic tumor. MR = magnetic resonance

Fig. 2. The excellent soft tissue contrast resolution of MR imaging showing invasion of adenocarcinoma into adjacent structures.
An axial contrast-enhanced computed tomography image (A) shows a heterogeneously enhancing mass exhibiting local invasion into the adjacent right brachiocephalic vein (arrowheads) in the right upper lobe. An axial T1-weighted image (B) shows abutment and the loss of fat plane between the mass and the right brachiocephalic vein (arrowheads), indicating that local invasion has occurred. Transbronchial biopsy was performed, and the histological diagnosis was an adenocarcinoma.
a significant finding because it implies that the lesion is not simply a benign cyst. Image subtraction is considered beneficial to evaluate enhancement (3).

Microscopic fat is easily detected on chemical shift imaging, a technique in which in-phase and out-of-phase images are obtained. If the signal intensity decreases significantly on the out-of-phase image, the lesion is likely to contain microscopic fat. Different fat-suppression techniques, such as fat saturation imaging, can be utilized to identify macroscopic or gross fat. This tissue shows a high signal intensity on T1- and T2-weighted images and again shows a signal decrease on fat-suppression sequences (6).

Diffusion-weighted MRI can provide useful information on tissue cellularity. The b-values of these images represent the degree of diffusion weighting: images with lower b values (50–100 s/mm²) are less affected by diffusion than those with higher b-values (800–1000 s/mm²). In high-cellularity tissues such as tumors, water diffusion is restricted, resulting in a higher signal intensity on higher b-value imaging (7). An apparent diffusion coefficient (ADC) map can be produced by reconstructing diffusion-weighted images. Malignant tumors have a low mean value on the ADC map (3, 8).

MRI has excellent soft tissue resolution; this aids in the evaluation of the origin and extent of lesions and their invasion into adjacent structures (Fig. 1). Abutment and the loss of fat plane are important MRI features to consider in the evaluation of local invasiveness (Fig. 2). Adherence to adjacent structures can be evaluated with cine imaging such as balanced steady-state free precession (3, 9).

The Korean Society of Thoracic Radiology recently proposed a standard mediastinal MRI protocol, which is summarized in Table 1.

### Prevascular Compartment

The ITMIG has defined the prevascular compartment of the mediastinum as the space between the posterior surface of the sternum and the anterior surface of the pericardium (Fig. 3). Approximately half of all mediastinal masses are located in the prevascular compartment, including a heterogeneous and diverse group of neoplasms (10). The prevalence of different prevascular mediastinal masses varies according to age and sex (11) (Table 2).

**Table 1. Standard Mediastinal MRI Protocol of the Korean Society of Thoracic Radiology**

<table>
<thead>
<tr>
<th>Contrast agent</th>
<th>Administer a single dose of contrast medium at 2 mL/s</th>
</tr>
</thead>
<tbody>
<tr>
<td>Preliminary imaging</td>
<td>Obtain variable scout images</td>
</tr>
<tr>
<td><strong>MRI Scan Order</strong></td>
<td><strong>Plane</strong></td>
</tr>
<tr>
<td>T2-weighted SSFSE</td>
<td>Axial and coronal</td>
</tr>
<tr>
<td>T2-weighted STIR</td>
<td>Axial</td>
</tr>
<tr>
<td>T1-weighted in- and out-of-phase</td>
<td>Axial</td>
</tr>
<tr>
<td>DW imaging (more than 3 b values)</td>
<td>Axial</td>
</tr>
<tr>
<td>T1-weighted fast-GRE</td>
<td>Axial and coronal or sagittal</td>
</tr>
<tr>
<td>Cine imaging (bSSFP)†</td>
<td>Coronal or sagittal</td>
</tr>
</tbody>
</table>

*Post-contrast images are obtained at 20–30 s, 60–70 s, 3 min, and 5 min. †Optional. bSSFP = balanced steady state free precession, DW = diffusion-weighted, ECG = electrocardiographic, GRE = gradient echo, MRI = magnetic resonance imaging, SSFSE = single shot fast spin echo, ST = slice thickness, STIR = short tau inversion recovery.
Table 2. Predominant Prevascular Mediastinal Masses according to Age and Sex

<table>
<thead>
<tr>
<th>Age (Years)</th>
<th>Benign Disease</th>
<th>Malignancy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Under 30s</td>
<td>Thymic hyperplasia or remnant (MC)</td>
<td>Aggressive lymphoma (MC)</td>
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<tr>
<td></td>
<td>Thymic bed cyst (2nd MC)</td>
<td>Malignant germ cell tumor</td>
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<tr>
<td></td>
<td>Benign teratoma</td>
<td></td>
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<tr>
<td>30s–40s</td>
<td>Thymic hyperplasia or remnant (MC)</td>
<td>Thymoma (MC)</td>
</tr>
<tr>
<td></td>
<td>Thymic bed cyst (2nd MC)</td>
<td>Aggressive lymphoma</td>
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<tr>
<td></td>
<td>Benign teratoma</td>
<td></td>
</tr>
<tr>
<td>40s–50s</td>
<td>Thymic bed cyst (MC)</td>
<td>Thymoma (MC)</td>
</tr>
<tr>
<td></td>
<td>Thymic hyperplasia or remnant (2nd MC)</td>
<td>Aggressive lymphoma</td>
</tr>
<tr>
<td></td>
<td>Benign teratoma</td>
<td></td>
</tr>
<tr>
<td>50s–60s</td>
<td>Thymic bed cyst</td>
<td>Thymoma (MC)</td>
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<tr>
<td></td>
<td></td>
<td>Thymic carcinoma</td>
</tr>
<tr>
<td>Over 60s</td>
<td>Thymic bed cyst</td>
<td>Thymoma (MC)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Thymic carcinoma</td>
</tr>
<tr>
<td>Female</td>
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<tr>
<td>Under 30s</td>
<td>Thymic hyperplasia or remnant</td>
<td>Aggressive lymphoma</td>
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<tr>
<td>30s–40s</td>
<td>Thymic hyperplasia or remnant (MC)</td>
<td>Aggressive lymphoma (MC)</td>
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<td></td>
<td>Thymic bed cyst (2nd MC)</td>
<td>Thymoma</td>
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<tr>
<td></td>
<td>Benign teratoma</td>
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<tr>
<td>40s–50s</td>
<td>Thymic bed cyst (MC)</td>
<td>Thymoma (MC)</td>
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<td></td>
<td>Thymic hyperplasia or remnant (2nd MC)</td>
<td>Thymic carcinoma</td>
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<td></td>
<td>Benign teratoma</td>
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<td>50s–60s</td>
<td>Thymic bed cyst</td>
<td>Thymoma (MC)</td>
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<td></td>
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<td>Thymic carcinoma</td>
</tr>
<tr>
<td>Over 60s</td>
<td>Thymic bed cyst (MC)</td>
<td>Thymoma (MC)</td>
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<tr>
<td></td>
<td>Benign teratoma</td>
<td>Thymic carcinoma</td>
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</tbody>
</table>

Both thymic cysts and bronchogenic cysts are classed as thymic bed cysts; thyroid goiter is excluded in this table. Carter et al. (12) reported that thyroid goiter and thymic malignancies are the most common causes of prevascular mediastinal masses in both male and female patients over 40 years old. MC = most common.

Fig. 4. A 40-year-old man with thymic hyperplasia (arrow).

A. An axial contrast-enhanced computed tomography image at the level of the ascending thoracic aorta shows a soft tissue lesion with a similar appearance to the normal thymus in the thymic bed of the prevascular mediastinum. B, C. On chemical shift imaging, the lesion shows a signal decrease on the out-of-phase image (C) compared to the in-phase image (B), supporting a diagnosis of thymic hyperplasia.
Thymic Hyperplasia
Thymic hyperplasia is observed as one of two characteristic patterns on imaging studies, depending on the patient’s age. In patients aged less than 40 years, it appears as a diffuse enlargement of thymic tissue compared with previous imaging studies, whereas in patients aged over 40 years, it presents as a soft tissue lesion similar in appearance to the normal thymus (1). Known causative factors include stress such as burns and injuries and previous histories of chemotherapy, radiation therapy, or corticosteroid treatment. Thymic hyperplasia is also associated with hyperthyroidism, myasthenia gravis, collagen vascular diseases, and human immunodeficiency virus (12).

Chemical shift MRI is considered beneficial in differentiating thymic hyperplasia from thymic tumors or other soft tissue malignancies. Thymic hyperplasia shows a signal decrease on out-of-phase imaging due to the presence of fat tissue (13, 14) (Fig. 4).

Thymic Cyst
When a cystic lesion is observed in the thymic bed of the prevascular mediastinum, it is likely to be a thymic cyst.

Fig. 6. A 52-year-old woman with a pericardial cyst in the right cardiophrenic angle. An axial fat-suppressed T2-weighted image shows fluid signal intensity within the mass.

Fig. 5. A 49-year-old woman with a thymic cyst (arrow).
A. An axial pre-contrast computed tomography image at the level of the ascending thoracic aorta shows an approximately 4-cm mass in the thymic bed of the prevascular mediastinum. The mass shows soft tissue attenuation (50 Hounsfield units). B, C. The mass shows high signal intensity on axial fat-suppressed T1-weighted (B) and T2-weighted (C) images. D. Subtraction imaging shows a lack of enhancement.
Thymic cysts can be either congenital or acquired. Acquired lesions may result from chemotherapy, radiation therapy, or thoracotomy (15, 16). Tomiyama et al. (17) reported that approximately three quarters of resected thymic cysts show soft tissue attenuation on pre-contrast CT. Similarly, Lee et al. (18) reported that the mean attenuation value of benign thymic bed cysts on pre-contrast CT was 31 Hounsfield units, a value higher than that of water. MRI is more useful than CT for evaluating thymic cysts because the cystic component can be more accurately assessed. Cysts typically show higher signal intensity than solid masses on T2-weighted images, but there is a significant overlap (19). Cysts show varying signal intensities on T1-weighted images. Confirming the lack of enhancement on post-contrast images is essential to avoid misinterpreting cysts as solid masses (19) (Fig. 5).

Fig. 7. A 63-year-old man with a thymoma (arrow).

A, B. An axial fat-suppressed T2-weighted image (A) shows hyperintensity within the mass, with inralesional foci of low signal intensity due to hemorrhage, flow void, or calcifications. The capsule is visible as a rim of low signal intensity on an axial fat-suppressed T1-weighted image (arrowheads, B). C, D. The mass has a high signal intensity on a higher b-value diffusion-weighted image (b = 800 s/mm², C). The apparent diffusion coefficient map (D) shows a low signal intensity, indicating diffusion restriction.
Pericardial Cyst

When cystic lesions are observed in a cardiophrenic angle, they are likely to be pericardial cysts. These are congenital anomalies, resulting from aberrations in the formation of coelomic (somatic) cavities (20). The right anterior cardiophrenic angle is their typical location (1). Similar to other mediastinal cysts, they show fluid signal intensity on MRI (Fig. 6). Occasionally, they are observed as high as the pericardial recess (21).

Thymoma

In adults, thymoma is the most common primary neoplasm in the prevascular mediastinum. Thymomas account for 20% of all mediastinal neoplasms (22). The Masaoka-Koga staging system is commonly used for thymomas. This includes four stages: stages I and II are early stages, while stages III and IV are advanced stages. Thymomas usually show low signal intensity on T1-weighted images and high signal intensity on T2-weighted images. They can show intralesional foci of low signal intensity on T2-weighted images.

Fig. 8. A 49-year-old man with a cystic thymoma.
A. An axial contrast-enhanced computed tomography image at the level of the ascending thoracic aorta shows an approximately 4.5-cm hypoattenuating mass with a thick wall in the thymic bed of the prevascular mediastinum. B. An axial fat-suppressed T2-weighted image shows a cystic area within the mass (white arrowheads). C. An axial post-contrast MR image shows an enhancing solid portion of the mass (black arrowheads). This patient also had pulmonary alveolar proteinosis in both lungs, which was confirmed by surgical biopsy.

Fig. 9. A 62-year-old woman with a mediastinal hemangioma.
A. An axial fat-suppressed T2-weighted image shows hyperintensity within the mass (arrow). B. An axial post-contrast MR image shows peripheral nodular enhancement (arrowhead).
images due to hemorrhage, flow void, or calcifications (23). Sakai et al. (24) reported that the presence of septa is a characteristic feature of thymoma, and this results in a lobulated shape. A lobulated or irregular border, necrotic or cystic changes, calcifications, lymphadenopathy, and the invasion of great vessels on CT or MRI have been identified as features indicative of high-risk thymoma or thymic carcinoma (22, 25, 26). Conversely, low-risk thymomas typically show a smooth border, an almost complete capsule, a septum, and homogeneous enhancement. The capsule is observed as a low signal intensity rim on MRI (27). Diffusion-weighted MRI is considered beneficial for identifying advanced stage thymomas. Abdel Razek et al. (8) reported that advanced stage thymomas show significantly lower mean ADC values than early stage thymomas, although there is some overlap (Fig. 7).

Thymomas can undergo degenerative cystic change. Cystic changes are typically limited to a focal area; however, extensive cystic change is sometimes observed (28). While these lesions may resemble other congenital cysts on CT

Fig. 10. A 21-year-old man with a T-cell lymphoblastic lymphoma. The mass has a lobular shape and a heterogeneous appearance. A. An axial fat-suppressed T2-weighted image shows hyperintensity within the mass. The encasement of adjacent vascular structures is also apparent. B. A coned-down axial fluorodeoxyglucose positron emission tomography/computed tomography image shows heterogeneous fluorodeoxyglucose uptake due to the presence of internal necrosis.

Fig. 11. A 35-year-old woman with a mature teratoma. A. An axial contrast-enhanced computed tomography image at the level of the left atrium shows a cystic mass in the left prevascular mediastinum, containing gross fat (asterisk), calcification (arrowhead), and regions of soft tissue (arrow). B, C. The gross fat (asterisk) has a high signal intensity on an axial T1-weighted image (B), whereas a signal decrease is observed on an axial fat-suppressed T2-weighted image (C).
or MRI, the presence of a solid portion, mural nodules, or septa within a cystic lesion in the prevascular mediastinum should prompt another diagnosis such as cystic thymoma (1, 29) (Fig. 8).

**Mediastinal Hemangioma**
Mediastinal hemangiomas only account for 0.5% of all mediastinal masses (30, 31). The majority arise in the prevascular mediastinum. MRI findings are similar to those in hepatic hemangioma. Lesions have low to intermediate signal intensity on T1-weighted images and high signal intensity on T2-weighted images. They can show three enhancement patterns on post-contrast images: uniform enhancement and peripheral nodular enhancement with or without centripetal enhancement (9, 32-34) (Fig. 9).

**Lymphoma**
Lymphoma should be considered in patients aged less than 50 years with a soft tissue mass in the prevascular mediastinum (11). A lobular shape and mild enhancement of the mass and the presence of conglomerate lymph nodes in the prevascular mediastinum should raise the suspicion of lymphoma. Lymphadenopathy elsewhere in the body also supports this diagnosis. The encasement of vascular structures without invasion is more often observed on CT or MRI scans of lymphoma than those of thymic epithelial tumors. Lymphomas show heterogeneous fluorodeoxyglucose uptake on fluorodeoxyglucose positron emission tomography/CT imaging due to internal necrosis (Fig. 10).

Considering these imaging findings and classic clinical “B” symptoms such as fever, night sweats, and weight loss, practitioners can confidently diagnose lymphoma (1).

**Mature Teratoma**
Mature teratomas are most frequently observed in young male patients aged less than 40 years (12). Teratomas comprise various internal constituents such as fat, soft tissue, calcification, and fluid. These are easily identifiable on MRI, with the exception of calcification, which is better assessed with CT. Intralesional fat (gross fat) usually has

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*Fig. 13. A 55-year-old man with an esophageal duplication cyst (asterisk) in the visceral mediastinum. The mass has a well-defined margin and an oval shape. On an axial fat-suppressed T2-weighted image, it can be observed that the mass has a high signal intensity and is attached to the intimal layer of the esophageal wall.*

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*Fig. 12. A 55-year-old woman with a bronchogenic cyst in the subcarinal area of the visceral mediastinum. A. A coronal fat-suppressed T2-weighted image shows hyperintensity within the mass. B. A fluid-fluid level (arrowheads) is apparent on an axial fat-suppressed T2-weighted image.*
high signal intensity on T1- and T2-weighted images, and the signal is decreased in fat-suppression sequences (35) (Fig. 11). The fat-fluid level is another diagnostic clue (36).

### Visceral Compartment

The visceral compartment is located between the pericardium and the visceral-paravertebral compartment line, an artificial line connecting each point 1 cm posterior to the ventral margin of the thoracic spine (Fig. 3). Most masses arising in this compartment are congenital cysts.

### Bronchogenic Cyst

The most common location of bronchogenic cysts is the visceral and paravertebral mediastinum near the tracheal carina, although these lesions can arise in any location (4). Patients are usually asymptomatic, although the mass effect on surrounding structures occasionally causes symptoms. Bronchogenic cysts have high signal intensity on T2-weighted images due to their high water content. Similar to other mediastinal cysts, they have variable signal intensities on T1-weighted images. Occasionally a fluid-fluid level is observed (20, 37) (Fig. 12).

### Esophageal Duplication Cyst

The esophagus is the second most common location in the gastrointestinal (GI) tract for duplication cysts. Intimate attachment to the GI tract, smooth muscle

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**Fig. 14.** A 57-year-old man with a mediastinal thyroid carcinoma originating from ectopic thyroid tissue.  
A. An axial T2-weighted image at the level of the thoracic inlet shows a septated cystic mass in the right upper visceral mediastinum with a fluid-fluid level (arrowheads) and a focal area of soft tissue (arrow). The histological diagnosis after surgical excision was a mediastinal thyroid carcinoma originating from ectopic thyroid tissue.  
B. The ectopic thyroid tissue (arrow) shows high attenuation and calcification on an axial pre-contrast computed tomography image.

**Fig. 15.** A 53-year-old man with a mediastinal liposarcoma.  
A. An axial contrast-enhanced computed tomography image at the level of the left ventricle shows a large heterogeneously enhancing mass in the visceral mediastinum with intralosomal gross fat (arrow). The mass compresses and displaces the heart and esophagus.  
B, C. On an axial T1-weighted image (B), the gross fat (arrow) has a high signal intensity, whereas on an axial post-contrast MR image (C), the gross fat (arrow) shows a signal decrease, and the mass exhibits heterogeneous enhancement.
within the wall, and an epithelial lining are the diagnostic
criteria for duplication cysts. These cysts usually have a
well-defined margin, an oval or tubular shape, and cystic
features. On MRI, they show high signal intensity on T2-
weighted images and variable signal intensities on T1-
weighted images (38, 39) (Fig. 13).

**Mediastinal Thyroid Carcinoma Originating from Ectopic
Thyroid Tissue**

The mediastinum is a rare location for ectopic thyroid
tissue. Diagnosing mediastinal ectopic thyroid tissue is
extremely difficult due to the presence of several differential
diagnoses such as thymic epithelial tumors and lymphoma.
Pre-contrast CT is useful in these cases considering that
ectopic thyroid tissue shows high attenuation (70 ± 10
Hounsfield units) compared with the skeletal muscle due to
its high iodine content (40). Furthermore, this tissue can
be evaluated with radionuclide imaging using technetium-
99m pertechnetate, iodine-123, or iodine-131, as it shows
radiiodine uptake (41). While malignant transformation of
ectopic thyroid tissue is exceedingly rare, the tissue should
be surgically resected to guard against this possibility (42,
43) (Fig. 14).

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**Fig. 16. A 79-year-old man with a recent history of pancreatitis who developed a mediastinal pancreatic pseudocyst.**

An axial fat-suppressed T2-weighted image (A) shows a loculated cystic lesion (asterisk) in the visceral mediastinum. This lesion shows communication (arrow) with the abdominal component on a coronal post-contrast MR image (B).

**Fig. 17. A 56-year-old man with a schwannoma in the right paravertebral mediastinum.** An axial fat-suppressed T2-weighted image shows central hypointense foci within the mass; these represent thickened fascicles (fascicular sign, arrowheads). The mass abuts the adjacent spinal nerve (arrow).

**Fig. 18. A 42-year-old man with a cystic schwannoma in the right paravertebral mediastinum.** An axial T2-weighted image shows the cystic nature of the mass and the thick appearance of its wall. A fluid-fluid level (arrowheads) is also observed within the mass.
Mediastinal Liposarcoma

Primary intrathoracic liposarcomas are rare, accounting for only 2.7% of liposarcomas (44). Mediastinal liposarcomas comprise only 0.1–0.8% of all mediastinal tumors (45). Differentiating liposarcoma from lipoma is relatively simple. In contrast to lipomas, liposarcomas usually have a heterogeneous appearance and show an internal area of soft tissue enhancement on MRI. The intralosomal fat (gross fat) in these lesions has a high signal intensity on T1- and T2-weighted images. This signal is decreased on fat-suppression images (46, 47) (Fig. 15).

Mediastinal Pancreatic Pseudocyst

Pancreatic pseudocysts are a complication of pancreatitis. When the pancreatic duct ruptures, proteolytic enzymes can affect the diaphragm, and the pseudocyst can extend to the intrathoracic mediastinum (48, 49). Thus, patients presenting with a loculated cystic lesion in the visceral mediastinum and a recent history of pancreatitis have a high risk of a pancreatic pseudocyst (50). These lesions have a cystic nature on MRI (20). Communication with their abdominal component is frequently observed (51) (Fig. 16).

Paravertebral Compartment

The paravertebral compartment is located between the visceral-paravertebral compartment line and the posterolateral aspect of the transverse processes of the thoracic spine (Fig. 3).

Neurogenic Tumors

As the paravertebral compartment includes the central canal and intervertebral foramen, which contain neural tissue, neurogenic tumors are the most commonly encountered masses in this compartment. MRI is useful to assess intraspinal extension of lesions through the neural foramen (1). Neurogenic tumors are typically divided into two groups depending upon their origin: first, neurogenic neoplasms of autonomic ganglia such as neuroblastomas and, second, nerve sheath tumors such as schwannomas and neurofibromas (52).

Schwannomas and neurofibromas are benign neoplasms that usually affect adolescents and adults aged less than 40 years (52). These nerve sheath tumors have certain characteristic MRI features. When multiple small hypointense foci with a ring-like shape are visible in a hyperintense mass on T2-weighted images, these indicate fascicular bundles and are termed fascicular signs. These are commonly observed in schwannomas (53) (Fig. 17). Schwannomas can undergo cystic degeneration, which is visualized as an area of very high signal intensity on T2-weighted images. A post-contrast image can help to differentiate between a schwannoma exhibiting cystic degeneration and other mediastinal cysts. Areas of cystic degeneration are typically associated with a well-defined margin and heterogeneous enhancement (Fig. 20).
location of primary thoracic neuroblastomas. These tumors typically affect infants and children aged less than 3 years. MRI can aid in the management of these patients as the images show the spinal canal in detail (55, 56) (Fig. 20).

Meningocele

When the leptomeninges herniate into an intervertebral foramen or a defect in a vertebral body, this causes the development of an anomalous paravertebral cystic mass termed as meningocele. Most meningoceles are detected in middle-aged adults (57). On MRI, meningoceles usually appear as paravertebral masses with a signal intensity equivalent to that of fluid. Widening of the intervertebral foramina can also be observed. Occasionally, neurogenic tumors with cystic changes resemble meningoceles. If a mass communicates with the thecal sac, it is more likely to be a meningocele (1, 20) (Fig. 21).

Plasmacytoma

A plasmacytoma is a malignant neoplasm of plasma cells and is thus a counterpart of multiple myeloma. Plasmacytomas are typically single lesions. There are two types of these tumors: osseous plasmacytomas and extramedullary plasmacytomas. Osseous plasmacytomas appear as osteolytic lesions with a bulky soft tissue component on MRI. They usually show iso-signal intensity on T2-weighted images. Large masses may show internal necrosis and heterogeneous enhancement on post-contrast images (58, 59) (Fig. 22).
CONCLUSION

MRI has become a valuable tool to evaluate mediastinal masses. In particular, MRI is useful for differentiating cystic from solid lesions, evaluating the invasion of a mass into adjacent structures, and characterizing tissue components such as fat or hemorrhage. MRI scans of the mediastinum, in comparison to other anatomical locations, are relatively simple to interpret. It is important to determine which of the three compartments the lesion is located. Fundamental knowledge about MRI findings and epidemiology can help radiologists and clinicians improve patient management (Table 3).

Conflicts of Interest
The authors have no potential conflicts of interest to disclose.

Table 3. Mediastinal Masses and Their Characteristic MRI Findings

<table>
<thead>
<tr>
<th>Lesion</th>
<th>MRI Findings</th>
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<tbody>
<tr>
<td>Prevascular compartment</td>
<td></td>
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<tr>
<td>Thymic hyperplasia</td>
<td>Signal decrease on out-of-phase image</td>
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<tr>
<td>Thymic cyst</td>
<td>Cystic lesion in the thymic bed; lack of enhancement</td>
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<tr>
<td>Bronchogenic cyst</td>
<td>Cystic lesion; fluid-fluid level</td>
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<tr>
<td>Pericardial cyst</td>
<td>Cystic lesion in the cardiophrenic angle</td>
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<td>Thymoma</td>
<td>Soft tissue mass in the thymic bed</td>
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<tr>
<td>Cystic thymoma</td>
<td>Soft tissue mass with focal cystic change in the thymic bed</td>
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<td>Mediastinal hemangioma</td>
<td>Soft tissue mass with a similar enhancement pattern to that of hepatic hemangioma</td>
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<tr>
<td>Lymphoma</td>
<td>Soft tissue mass with a heterogeneous appearance; lymphadenopathy</td>
</tr>
<tr>
<td>Mature teratoma</td>
<td>A variety of internal contents (fat, soft tissue, calcification, and fluid)</td>
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<td>Visceral compartment</td>
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<tr>
<td>Bronchogenic cyst</td>
<td>Cystic lesion (near the tracheal carina); fluid-fluid level</td>
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<td>Esophageal duplication cyst</td>
<td>Cystic lesion with intimal esophageal attachment</td>
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<tr>
<td>Mediastinal thyroid carcinoma</td>
<td>Cystic lesion with an internal soft tissue area</td>
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<tr>
<td>Mediastinal liposarcoma</td>
<td>Soft tissue mass with intralesional fat, a heterogeneous appearance, and an internal enhancing soft tissue area</td>
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<tr>
<td>Mediastinal pancreatic pseudocyst</td>
<td>Loculated cystic lesion communicating with the abdominal component</td>
</tr>
<tr>
<td>Paravertebral compartment</td>
<td></td>
</tr>
<tr>
<td>Schwannoma</td>
<td>Multiple hypointense, small, ring-like structures corresponding to fascicular bundles (fascicular sign)</td>
</tr>
<tr>
<td>Cystic schwannoma</td>
<td>Cystic mass with peripheral enhancement and a thick and irregular wall</td>
</tr>
<tr>
<td>Neurofibroma</td>
<td>Central hypointensity and higher peripheral intensity (target sign)</td>
</tr>
<tr>
<td>Neuroblastoma</td>
<td>Soft tissue mass; patients younger than three years old</td>
</tr>
<tr>
<td>Meningocele</td>
<td>Cystic lesion communicating with the thecal sac</td>
</tr>
<tr>
<td>Plasmacytoma</td>
<td>Iso-signal intensity on T2-weighted images</td>
</tr>
</tbody>
</table>

Cystic lesions show high signal intensity on T2-weighted images, and sometimes show high or intermediate signal intensity on T1-weighted images.

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