Fleischner Society: Glossary of Terms for Thoracic Imaging

Members of the Fleischner Society compiled a glossary of terms for thoracic imaging that replaces previous glossaries published in 1984 and 1996 for thoracic radiography and computed tomography (CT), respectively. The need to update the previous versions came from the recognition that new words have emerged, others have become obsolete, and the meaning of some terms has changed. Brief descriptions of some diseases are included, and pictorial examples (chest radiographs and CT scans) are provided for the majority of terms.

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The present glossary is the third prepared by members of the Fleischner Society and replaces the glossaries of terms for thoracic radiology (1) and CT (2), respectively. The impetus to combine and update the previous versions came from the recognition that with the recent developments in imaging new words have arrived, others have become obsolete, and the meaning of some terms has changed. The intention of this latest glossary is not to be exhaustive but to concentrate on those terms whose meaning may be problematic. Terms and techniques not used exclusively in thoracic imaging are not included.

Two new features are the inclusion of brief descriptions of the idiopathic interstitial pneumonias (IIPs) and pictorial examples (chest radiographs and computed tomographic [CT] scans) for the majority of terms. The decision to include vignettes of the IIPs (but not other pathologic entities) was based on the perception that, despite the recent scrutiny and reclassification, the IIPs remain a confusing group of diseases. We trust that the illustrations enhance, but do not distract from, the definitions. In this context, the figures should be regarded as of less importance than the text—they are merely examples and should not be taken as representing the full range of possible imaging appearances (which may be found in the references provided in this glossary or in comprehensive textbooks).

We hope that this glossary of terms will be helpful, and it is presented in the spirit of the sentiment of Edward J. Huth that “scientific writing calls for precision as much in naming things and concepts as in presenting data” (3). It is right to repeat the request with which the last Fleischner Society glossary closed: “[U]se of words is inherently controversial and we are pleased to invite readers to offer improvements to our definitions” (2).

**Glossary**

**acinus**

*Anatomy.*—The acinus is a structural unit of the lung distal to a terminal bronchiole and is supplied by first-order respiratory bronchioles; it contains alveolar ducts and alveoli. It is the largest unit in which all airways participate in gas exchange and is approximately 6–10 mm in diameter. One secondary pulmonary lobe contains between three and 25 acini (4).

*Radiographs and CT scans.*—Individual normal acini are not visible, but acinar arteries can occasionally be identified on thin-section CT scans. Accumulation of pathologic material in acini may be seen as poorly defined nodular opacities on chest radiographs and thin-section CT images. (See also nodules.)

**acute interstitial pneumonia, or AIP**

*Pathology.*—The term *acute interstitial pneumonia* is reserved for diffuse alveolar damage of unknown cause. The acute phase is characterized by edema and hyaline membrane formation. The later phase is characterized by airspace and/or interstitial organization (5). The histologic pattern is indistinguishable from that of acute respiratory distress syndrome.

*Radiographs and CT scans.*—In the acute phase, patchy bilateral ground-glass opacities are seen (6), often with some spurring of individual lobules, producing a geographic appearance; dense opacification is seen in the dependent lung (Fig 1). In the organizing phase, architectural distortion, traction bronchiectasis, cysts, and reticular opacities are seen (7).

**air bronchogram**

*Radiographs and CT scans.*—An air bronchogram is a pattern of air-filled (low-attenuation) bronchi on a background of opaque (high-attenuation) airless lung (Fig 2). The sign implies (a) patency of proximal airways and (b) evacuation of alveolar air by means of absorption (atelectasis) or replacement (eg, pneumonia) or a combination of these processes. In rare cases, the displacement of air is the result of marked interstitial expansion (eg, lymphoma) (8).
**air crescent**

*Radiographs and CT scans.*—An air crescent is a collection of air in a crescentic shape that separates the wall of a cavity from an inner mass (Fig 3). The air crescent sign is often considered characteristic of either *Aspergillus* colonization of preexisting cavities or retraction of infarcted lung in angioinvasive aspergillosis (9,10). However, the air crescent sign has also been reported in other conditions, including tuberculosis, Wegener granulomatosis, intracavitary hemorrhage, and lung cancer. (See also *mycetoma.*)

**air trapping**

*Pathophysiology.*—Air trapping is retention of air in the lung distal to an obstruction (usually partial).

**airspace**

*Anatomy.*—An airspace is the gas-containing part of the lung, including the respiratory bronchioles but excluding purely conducting airways, such as terminal bronchioles.

*Radiographs and CT scans.*—This term is used in conjunction with consolidation, opacity, and nodules to designate the filling of airspaces with the products of disease (14).

**aortopulmonary window**

*Anatomy.*—The aortopulmonary window is the mediastinal region bounded anteriorly by the ascending aorta, posteriorly by the descending aorta, cranially by the aortic arch, inferiorly by the left pulmonary artery, medially by the ligamentum arteriosum, and laterally by the pleura and left lung (15,16).

*Radiographs and CT scans.*—Focal concavity in the left mediastinal border below the aorta and above the left pulmonary artery can be seen on a frontal radiograph (Fig 5). Its appearance may be modified by tortuosity of the aorta. The aortopulmonary window is a common site of lymphadenopathy in a variety of inflammatory and neoplastic diseases.

**apical cap**

*Pathology.*—An apical cap is a caplike lesion at the lung apex, usually caused by intrapulmonary and pleural fibrosis pulling down extrapleural fat (17) or possibly by chronic ischemia resulting in hyaline plaque formation on the visceral pleura (18). The prevalence increases with age. It can also be seen in hematoma resulting from aortic rupture or in other fluid collection associated with infection or tumor, either outside the parietal pleura or loculated within the pleural space (19).

*Radiographs and CT scans.*—The usual appearance is of homogeneous soft-tissue attenuation capping the ex-
treme lung apex (uni- or bilaterally), with a sharp or irregular lower border (Fig 6). Thickness is variable, ranging up to about 30 mm (17). An apical cap occasionally mimics apical consolidation on transverse CT scans.

**architectural distortion**

Pathology.—Architectural distortion is characterized by abnormal displacement of bronchi, vessels, fissures, or septa caused by diffuse or localized lung disease, particularly interstitial fibrosis.

CT scans.—Lung anatomy has a distorted appearance and is usually associated with pulmonary fibrosis (Fig 7) and accompanied by volume loss.

**atelectasis**

Pathophysiology.—Atelectasis is reduced inflation of all or part of the lung (20). One of the commonest mechanisms is resorption of air distal to airway obstruction (eg, an endobronchial neoplasm) (21). The synonym collapse is often used interchangeably with atelectasis, particularly when it is severe or accompanied by obvious increase in lung opacity.

Radiographs and CT scans.—Reduced volume is seen, accompanied by increased opacity (chest radiograph) or attenuation (CT scan) in the affected part of the lung (Fig 8). Atelectasis is often associated with abnormal displacement of fissures, bronchi, vessels, diaphragm, heart, or mediastinum (22). The distribution can be lobar, segmental, or subsegmental. Atelectasis is often qualified by descriptors such as linear, discoid, or platelike. (See also linear atelectasis, rounded atelectasis.)

**azygos fissure**

See fissure.

**azygoesophageal recess**

Anatomy.—The azygoesophageal recess is a right posterior mediastinal recess into which the edge of the right lower lobe extends. It is limited superiority by the azygos arch, posteriorly by the azygos vein and pleura anterior to the vertebral column, and medially by the esophagus and adjacent structures.

Radiographs and CT scans.—On a frontal chest radiograph, the recess is seen as a vertically oriented interface between the right lower lobe and the adjacent mediastinum (the medial limit of the recess). Superiorly, the interface is seen as a smooth arc with convexity to the left. Disappearance or distortion of part of the interface suggests disease (eg, subcarinal lymphadenopathy). On CT scans, the recess (Fig 9) merits attention because small lesions located in the recess will often be invisible on chest radiographs (23).

**beaded septum sign**

CT scans.—This sign consists of irregular and nodular thickening of interlobular septa reminiscent of a row of beads (Fig 10). It is frequently seen in lymphangitic spread of cancer and less often in sarcoidosis (24).

**bleb**

Anatomy.—A bleb is a small gas-containing space within the visceral pleura or in the subpleural lung, not larger than 1 cm in diameter (25).

CT scans.—A bleb appears as a thin-walled cystic air space contiguous with the pleura. Because the arbitrary (size) distinction between a bleb and
bulla is of little clinical importance, the use of this term by radiologists is discouraged.

**Bronchiectasis**

**Pathology.**—Bronchiectasis is irreversible localized or diffuse bronchial dilatation, usually resulting from chronic infection, proximal airway obstruction, or congenital bronchial abnormality (26). (See also traction bronchiectasis.)

**Radiographs and CT scans.**—Morphologic criteria on thin-section CT scans include bronchial dilatation with respect to the accompanying pulmonary artery (signet ring sign), lack of tapering of bronchi, and identification of bronchi within 1 cm of the pleural surface (27) (Fig 11). Bronchiectasis may be classified as cylindrical, varicose, or cystic, depending on the appearance of the affected bronchi. It is often accompanied by bronchial wall thickening, mucoid impaction, and small-airways abnormalities (27–29). (See also signet ring sign.)

**Bronchiole**

**Anatomy.**—Bronchioles are non–cartilage-containing airways. Terminal bronchioles are the most distal of the purely conducting airways; they give rise to respiratory bronchioles, from which the alveoli arise and permit gas exchange. Respiratory bronchioles branch into multiple alveolar ducts (30).

**Bronchocele**

**Pathology.**—A bronchocele is bronchial dilatation due to retained secretions (mucoid impaction) usually caused by proximal obstruction, either congenital (eg, bronchial atresia) or acquired (eg, obstructing cancer) (34).

**Radiographs and CT scans.**—A bronchocele is a tubular or branching Y- or V-shaped structure that may resemble a gloved finger (Fig 13). The CT attenuation of the mucus is generally that of soft tissue but may be modified by its composition (eg, high-attenuation material in allergic bronchopulmonary aspergillosis). In the case of bronchial atresia, the surrounding lung may be of decreased attenuation because of reduced ventilation and, thus, perfusion.
bronchocentric

CT scans.—This descriptor is applied to disease that is conspicuously centered on macroscopic bronchovascular bundles (Fig 14). Examples of diseases with a bronchocentric distribution include sarcoidosis (35), Kaposi sarcoma (36), and organizing pneumonia (37).

broncholith

Pathology.—A broncholith, a calcified peribronchial lymph node that erodes into an adjacent bronchus, is most often the consequence of Histoplasma or tuberculous infection.

Radiographs and CT scans.—The imaging appearance is of a small calcific focus in or immediately adjacent to an airway (Fig 15), most frequently the right middle lobe bronchus. Broncholiths are readily identified on CT scans (38). Distal obstructive changes may include atelectasis, mucoid impaction, and bronchiectasis.

bulla

Pathology.—An airspace measuring more than 1 cm—usually several centimeters—in diameter, sharply demarcated by a thin wall that is no greater than 1 mm in thickness. A bulla is usually accompanied by emphysematous changes in the adjacent lung. (See also bullous emphysema.)

Radiographs and CT scans.—A bulla appears as a rounded focal lucency or area of decreased attenuation, 1 cm or more in diameter, bounded by a thin wall (Fig 16). Multiple bullae are often present and are associated with other signs of pulmonary emphysema (centrilobular and paraseptal).

cavity

Radiographs and CT scans.—A cavity is a gas-filled space, seen as a lucency or low-attenuation area, within pulmonary consolidation, a mass, or a nodule (Fig 17). In the case of cavitating consolidation, the original consolidation may resolve and leave only a thin wall. A cavity is usually produced by the expulsion or drainage of a necrotic part of the lesion via the bronchial tree. It sometimes contains a fluid level. Cavity is not a synonym for abscess.

centrilobular

Anatomy.—Centrilobular describes the region of the bronchiolovascular core of a secondary pulmonary lobule (4,39,40). This term is also used by pathologists to describe the location of lesions beyond the terminal bronchiole that center on respiratory bronchioles or even alveolar ducts.

CT scans.—A small dotlike or linear opacity in the center of a normal secondary pulmonary lobule, most obvious within 1 cm of a pleural surface, represents the intralobular artery (approximately 1 mm in diameter) (41). Centrilobular abnormalities include (a) nodules, (b) a tree-in-bud pattern indicating small-airways disease, (c) increased vis-
ibility of centrilobular structures due to thickening or infiltration of the adjacent interstitium, or (d) abnormal areas of low attenuation caused by centrilobular emphysema (4). (See also lobular core structures.)

**centrilobular emphysema**

*Pathology.*—Centrilobular emphysema is characterized by destroyed centrilobular alveolar walls and enlargement of respiratory bronchioles and associated alveoli (42,43). This is the commonest form of emphysema in cigarette smokers.

*CT scans.*—CT findings are centrilobular areas of decreased attenuation, usually without visible walls, of nonuniform distribution and predominantly located in upper lung zones (44) (Fig 18). The term **centriacinar emphysema** is synonymous. (See also emphysema.)

**collapse**

See atelectasis.

**consolidation**

*Pathology.*—Consolidation refers to an exudate or other product of disease that replaces alveolar air, rendering the lung solid (as in infective pneumonia).

*Radiographs and CT scans.*—Consolidation appears as a homogeneous increase in pulmonary parenchymal attenuation that obscures the margins of vessels and airway walls (45) (Fig 19). An air bronchogram may be present. The attenuation characteristics of consolidated lung are only rarely helpful in differential diagnosis (eg, decreased attenuation in lipoid pneumonia [46] and increased in amiodarone toxicity [47]).

**crazy-paving pattern**

*CT scans.*—This pattern appears as thickened interlobular septa and intralobular lines superimposed on a background of ground-glass opacity (Fig 20), resembling irregularly shaped paving stones. The crazy-paving pattern is often sharply demarcated from more normal lung and may have a geographic outline. It was originally reported in patients with alveolar proteinosis (48) and is also encountered in other diffuse lung diseases (49) that affect both the interstitial and airspace compartments, such as lipoid pneumonia (50).

**cyst**

*Pathology.*—A cyst is any round circumscribed space that is surrounded by an epithelial or fibrous wall of variable thickness (51).

*Radiographs and CT scans.*—A cyst appears as a round parenchymal lucency or low-attenuating area with a well-defined interface with normal lung. Cysts have variable wall thickness but are usually thin-walled (<2 mm) and occur without associated pulmonary emphysema (Fig 21). Cysts in the lung usually contain air but occasionally contain fluid or solid material. The term is often used to describe enlarged thin-walled airspaces in patients with lymphangioleiomyomatosis (52) or Langerhans cell histiocytosis (53); thicker-walled honeycomb cysts are seen in patients with end-stage fibrosis (54). (See also bleb, bulla, honeycombing, pneumatocele.)

**cryptogenic organizing pneumonia, or COP**

See organizing pneumonia.
desquamative interstitial pneumonia, or DIP
Pathology.—Histologically, DIP is characterized by the widespread accumulation of an excess of macrophages in the distal airspaces. The macrophages are uniformly distributed, unlike in respiratory bronchiolitis–interstitial lung disease, in which the disease is conspicuously bronchiocentric. Interstitial involvement is minimal. Most cases of DIP are related to cigarette smoking, but a few are idiopathic or associated with rare inborn errors of metabolism (5).

Radiographs and CT scans.—Ground-glass opacity is the dominant abnormality and tends to have a basal and peripheral distribution (Fig 22). Microcystic or honeycomb changes in the area of ground-glass opacity are seen in some cases (55).

diffuse alveolar damage, or DAD
See acute interstitial pneumonia.

emphysema
Pathology.—Emphysema is characterized by permanently enlarged airspaces distal to the terminal bronchiole with destruction of alveolar walls (42,43). Absence of “obvious fibrosis” was historically regarded as an additional criterion (42), but the validity of that criterion has been questioned because some interstitial fibrosis may be present in emphysema secondary to cigarette smoking (56,57). Emphysema is usually classified in terms of the part of the acinus predominantly affected: proximal (centriacinar, more commonly termed centrilobular, emphysema), distal (paraseptal emphysema), or whole acinus (panacinar or, less commonly, panlobular emphysema).

CT scans.—The CT appearance of emphysema consists of focal areas or regions of low attenuation, usually without visible walls (58). In the case of panacinar emphysema, decreased attenuation is more diffuse. (See also bullous emphysema, centrilobular emphysema, panacinar emphysema, paraseptal emphysema.)

fissure
Anatomy.—A fissure is the infolding of visceral pleura that separates one lobe or part of a lobe from another; thus, the interlobar fissures are produced by two layers of visceral pleura. Supernumerary fissures usually separate segments rather than lobes. The azygos fissure, unlike the other fissures, is formed by two layers each of visceral and parietal pleura. All fissures (apart from the azygos fissure) may be incomplete.

Radiographs and CT scans.—Fissures appear as linear opacities, normally 1 mm or less in thickness, that correspond in position and extent to the anatomic fissural separation of pulmonary lobes or segments. Qualifiers include minor, major, horizontal, oblique, accessory, anomalous, azygos, and inferior accessory.

folded lung
See rounded atelectasis.

fungus ball
See mycetoma.

gas trapping
See air trapping.

ground-glass nodule
See nodule.

ground-glass opacity
Radiographs and CT scans.—On chest radiographs, ground-glass opacity appears as an area of hazy increased lung opacity, usually extensive, within which margins of pulmonary vessels may be indistinct. On CT scans, it appears as hazy increased opacity of lung, with preservation of bronchial and vascular margins (Fig 23). It is caused by partial filling of airspaces, interstitial thickening (due to fluid, cells, and/or fibrosis), partial collapse of alveoli, increased capillary blood volume, or a combination of these, the common factor being the partial displacement of air (59,60). Ground-glass opacity is less opaque than consolidation, in which bronchovascular margins are obscured. (See also consolidation.)
hypothesis sign

*CT scans.*—The halo sign is a CT finding of ground-glass opacity surrounding a nodule or mass (Fig 24). It was first described as a sign of hemorrhage around foci of invasive aspergillosis (61). The halo sign is nonspecific and may also be caused by hemorrhage associated with other types of nodules (62) or by local pulmonary infiltration by neoplasm (eg, adenocarcinoma). (See also reversed halo sign.)

**hilum**

*Anatomy.*—*Hilum* is a generic term that describes the indentation in the surface of an organ, where vessels and nerves connect with the organ. It is the site on the medial aspect of the lung where the vessels and bronchi enter and leave the lung.

*Radiographs and CT scans.*—A hilum appears as a composite opacity at the root of each lung produced by bronchi, arteries, veins, lymph nodes, nerves, and other tissue. The terms *hilum* (singular) and *hila* (plural) are preferred to *hilus* and *hili* respectively; the adjectival form is *hilar.*

**honeycombing**

*Pathology.*—Honeycombing represents destroyed and fibrotic lung tissue containing numerous cystic airspaces with thick fibrous walls, representing the late stage of various lung diseases, with complete loss of acinar architecture. The cysts range in size from a few millimeters to several centimeters in diameter, have variable wall thickness, and are lined by metaplastic bronchiolar epithelium (51).

*Radiographs and CT scans.*—On chest radiographs, honeycombing appears as closely approximated ring shadows, typically 3–10 mm in diameter with walls 1–3 mm in thickness, that resemble a honeycomb; the finding implies end-stage lung disease. On CT scans, the appearance is of clustered cystic air spaces, typically of comparable diameters on the order of 3–10 mm but occasionally as large as 2.5 cm (Fig 25). Honeycombing is usually subpleural and is characterized by well-defined walls (54). It is a CT feature of established pulmonary fibrosis (5). Because honeycombing is often considered specific for pulmonary fibrosis and is an important criterion in the diagnosis of usual interstitial pneumonia (63), the term should be used with care, as it may directly impact patient care.

**idiopathic pulmonary fibrosis**

*Pathology.*—Idiopathic pulmonary fibrosis is a specific form of chronic fibrosing interstitial pneumonia of unknown cause and is characterized by a histologic pattern of usual interstitial pneumonia (5,64).

*Radiographs and CT scans.*—The typical imaging findings are reticular opacities and honeycombing, with a predominantly peripheral and basal distribution (Fig 26). Ground-glass opacity, if present, is less extensive than reticular and honeycombing patterns. The typical radiologic findings (65,66) are also encountered in usual interstitial pneumonia secondary to specific causes, such as asbestos-induced pulmonary fibrosis (asbestosis), and the diagnosis is usually one of exclusion. (See also usual interstitial pneumonia.)
infarction
Pathology.—Infarction is a process that may result in ischemic necrosis, usually the consequence of vascular compromise such as occlusion of a feeding pulmonary artery by an embolus (venous infarction is rare but recognized). Necrosis is relatively uncommon because tissue viability is maintained by the bronchial arterial blood supply. Pulmonary infarction may be secondary to a vasculitis (eg, Wegener granulomatosis).

Radiographs and CT scans.—A pulmonary infarct is typically triangular or dome-shaped, with the base abutting the pleura and the apex directed toward the hilum (Fig 27). The opacity represents local hemorrhage with or without central tissue necrosis (67,68).

infiltrate
Radiographs and CT scans.—Formerly used as a term to describe a region of pulmonary opacification caused by air-space or interstitial disease seen on radiographs and CT scans. Infiltrate remains controversial because it means different things to different people (69). The term is no longer recommended, and has been largely replaced by other descriptors. The term opacity, with relevant qualifiers, is preferred.

interlobular septal thickening
Radiographs and CT scans.—This finding is seen on chest radiographs as thin linear opacities at right angles to and in contact with the lateral pleural surfaces near the lung bases (Kerley B lines); it is seen most frequently in lymphangitic spread of cancer or pulmonary edema. Kerley A lines are predominantly situated in the upper lobes, are 2–6 cm long, and can be seen as fine lines radially oriented toward the hila. In recent years, the anatomically descriptive terms septal lines and septal thickening have gained favor over Kerley lines. On CT scans, disease affecting one of the components of the septa (see interlobular septum) may be responsible for thickening and so render septa visible. On thin-section CT scans, septal thickening may be smooth or nodular (70) (Fig 28), which may help refine the differential diagnosis. (See also interlobular septum, beaded septum.)

interlobular septum
Anatomy.—Interlobular septa are sheet-like structures 10–20-mm long that form the borders of lobules; they are more or less perpendicular to the pleura in the periphery. Interlobular septa are composed of connective tissue and contain lymphatic vessels and pulmonary venules.

Radiographs and CT scans.—Interlobular septa appear as thin linear opacities between lobules (Fig 29); these septa are to be distinguished from centrilobular structures. They are not usually seen in the healthy lung (normal septa are approximately 0.1 mm thick) but are clearly visible when thickened (eg, by pulmonary edema). (See also interlobular septal thickening, lobule.)
**interstitial emphysema**

**Pathology.**—Interstitial emphysema is characterized by air dissecting within the interstitium of the lung, typically in the peribronchovascular sheaths, interlobular septa, and visceral pleura. It is most commonly seen in neonates receiving mechanical ventilation.

**Radiographs and CT scans.**—Interstitial emphysema is rarely recognized radiographically in adults and is infrequently seen on CT scans (Fig 30). It appears as perivascular lucent or low-attenuating halos and small cysts (71,72).

**interstitium**

**Anatomy.**—The interstitium consists of a continuum of connective tissue throughout the lung comprising three subdivisions: (a) the bronchovascular (axial) interstitium, surrounding and supporting the bronchi, arteries, and veins from the hilum to the level of the respiratory bronchiole; (b) the parenchymal (acinar) interstitium, situated between alveolar and capillary basement membranes; and (c) the subpleural connective tissue contiguous with the interlobular septa (73).

**juxtaphrenic peak**

**Radiographs and CT scans.**—A juxtaphrenic peak is a small triangular opacity based at the apex of the dome of a hemidiaphragm, associated with upper lobe volume loss of any cause (eg, postirradiation fibrosis or upper lobectomy) (74). It is most readily appreciated on a frontal chest radiograph (Fig 32). The peak is caused by upward retraction of the inferior accessory fissure (75) or an intrapulmonary septum associated with the pulmonary ligament (76).

**intralobular lines**

**CT scans.**—Intralobular lines are visible as fine linear opacities in a lobule when the intralobular interstitial tissue is abnormally thickened (Fig 31). When numerous, they may appear as a fine reticular pattern. Intralobular lines may be seen in various conditions, including interstitial fibrosis and alveolar proteinosis (41).

**linear atelectasis**

**Radiographs and CT scans.**—Linear atelectasis is a focal area of subsegmental atelectasis with a linear configuration, almost always extending to the pleura (74). It is commonly horizontal but sometimes oblique or vertical. The thickness of the atelectasis may range from a few millimeters to more than 1 cm (Fig 33). Linear atelectasis is also referred to as discoid or platelike atelectasis. (See also atelectasis.)

**lobe**

**Anatomy.**—The lobe is the principal division of the lungs (normally, three lobes on the right and two on the left); each lobe is enveloped by visceral pleura, except at the lung root (hilum) and when an interlobar fissure is incomplete.
lobular core structures

Anatomy. — Lobular core structures are the central structures in secondary pulmonary lobules and consist of a centrilobular artery and bronchiole (40).

CT scans. — The pulmonary artery and its immediate branches are visible in the center of a secondary lobule on thin-section CT scans, particularly if thickened (eg, by pulmonary edema) (Fig 34). These arteries measure approximately 0.5–1.0 mm in diameter. The centrilobular structures, or core structures, include bronchioles and their accompanying pulmonary arterioles and lymphatic vessels. The connective-tissue septa surrounding the pulmonary lobule—the interlobular septa, which contain veins and lymphatic vessels—are best developed in the periphery in the anterior, lateral, and juxtamediastinal regions of the upper and middle lobes.

CT scans. — On thin-section CT scans, the three basic components of the lobule—the interlobular septa and septal structures, the central lobular region (centrilobular structures), and the lobular parenchyma—can be identified, particularly in disease states. Peripheral lobules are more uniform in appearance and pyramidal in shape than are central lobules (4) (Fig 35). (See also interlobular septa, lobular core structures.)

lymphadenopathy

Pathology. — By common usage, the term lymphadenopathy is usually restricted to enlargement, due to any cause, of the lymph nodes. Synonyms include lymph node enlargement (preferred) and adenopathy.

CT scans. — There is a wide range in the size of normal lymph nodes. Mediastinal and hilar lymph nodes range in size from sub-CT resolution to 12 mm. Somewhat arbitrary thresholds for the upper limit of normal of 1 cm in short-axis diameter for mediastinal nodes (79) and 3 mm for most hilar nodes (80) have been reported, but size criteria do not allow reliable differentiation between healthy and diseased lymph nodes (Fig 36).

lymphoid interstitial pneumonia, or LIP

Pathology. — LIP is a rare disease characterized by diffuse pulmonary lymphoid proliferation with predominant interstitial involvement. It is included in the spectrum of interstitial pneumonias and is distinct from diffuse lymphomas of the lung. Features include diffuse hyperplasia of bronchus-associated lymphoid tissue and diffuse polyclonal lym-
phoid cell infiltrates surrounding the airways and expanding the lung interstitium. LIP is usually associated with autoimmune diseases or human immunodeficiency virus infection (5,81).

CT scans.—Ground-glass opacity is the dominant abnormality, and thin-walled perivascular cysts may be present (Fig 37). Lung nodules, a reticular pattern, interlobular septal and bronchovascular thickening, and widespread consolidation may also occur (82,83).

mass

Radiographs and CT scans.—A mass is any pulmonary, pleural, or mediastinal lesion seen on chest radiographs as an opacity greater than 3 cm in diameter (without regard to contour, border, or density characteristics). Mass usually implies a solid or partly solid opacity. CT allows more exact evaluation of size, location, attenuation, and other features. (See also nodule.)

mediastinal compartments

Anatomy.—Nominal anatomic compartments of the mediastinum include the anterior, middle, posterior, and (in some schemes) superior compartments. The anterior compartment is bounded anteriorly by the sternum and posteriorly by the anterior surface of the pericardium, the ascending aorta, and the brachiocephalic vessels. The middle compartment is bounded by the posterior margin of the anterior division and the anterior margin of the posterior division. The posterior compartment is bounded anteriorly by the posterior margins of the pericardium and great vessels and posteriorly by the thoracic vertebral bodies. In the four-compartment model, the superior compartment is defined as the compartment above the plane between the sternal angle to the T4-5 intervertebral disk or, more simply, above the aortic arch (84,85). Exact anatomic boundaries between the compartments do not exist, and there are no barriers (other than the pericardium) to prevent the spread of disease between compartments. Other classifications exist, but the three- and four-compartment models are the most commonly used.

micronodule

CT scans.—A micronodule is a discrete, small, round, focal opacity. A variety of diameters have been used in the past to define a micronodule; for example, a diameter of no greater than 7 mm (86). Use of the term is most often limited to nodules with a diameter of less than 5 mm (87) or less than 3 mm (88). It is recommended that the term be reserved for opacities less than 3 mm in diameter. (See also nodule, miliary pattern.)

mosaic attenuation pattern

CT scans.—This pattern appears as patchwork of regions of differing attenuation that may represent (a) patchy interstitial disease, (b) obliterative small-airways disease (Fig 39), or (c) occlusive vascular disease (89). Mosaic attenuation pattern is a more inclusive term than the original terms mosaic oligemia and perfusion (90). Air trapping secondary to bronchial or bronchiolar obstruction may produce focal zones of decreased attenuation, an appearance that can be enhanced by using expiratory CT (91,92). The mosaic attenuation pattern can also be produced by interstitial lung disease characterized by ground-glass opacity; in this situation, areas of higher attenuation represent the interstitial process and areas of lower attenuation represent the normal lung.

mosaic oligemia, perfusion

See mosaic attenuation pattern.
mycetoma

Pathology.—A mycetoma is a discrete mass of intertwined hyphae, usually of an Aspergillus species, matted together by mucus, fibrin, and cellular debris colonizing a cavity, usually from prior fibrocavitary disease (eg, tuberculosis or sarcoidosis).

Radiographs and CT scans.—A mycetoma may move to a dependent location when the patient changes position and may show an air crescent sign (Fig 40). CT scans may show a spongelike pattern and foci of calcification in the mycetoma (93). A synonym is fungus ball. (See also air crescent.)

nodule

Radiographs and CT scans.—The chest radiographic appearance of a nodule is a rounded opacity, well or poorly defined, measuring up to 3 cm in diameter. (a) Acinar nodules are round or ovoid poorly defined pulmonary opacities approximately 5–8 mm in diameter, presumed to represent an anatomic acinus rendered opaque by consolidation. This classification is used only in the presence of numerous such opacities. (b) A pseudonodule mimics a pulmonary nodule; it represents, for example, a rib fracture, a skin lesion, a device on the surface of the patient, anatomic variants, or composite areas of increased opacity (94).

On CT scans, a nodule appears as a rounded or irregular opacity, well or poorly defined, measuring up to 3 cm in diameter (Fig 42). (a) Centrilobular nodules appear separated by several millimeters from the pleural surfaces, fissures, and interlobular septa. They may be of soft-tissue or ground-glass attenuation. Ranging in size from a few millimeters to a centimeter, centrilobular nodules are usually ill-defined (4). (b) A micronodule is less than 3 mm in diameter (see also micronodule). (c) A ground-glass nodule (synonym, nonsolid nodule) manifests as hazy increased attenuation in the lung that does not obliterate the bronchial and vascular margins. (d) A solid nodule has homogenous soft-tissue attenuation. (e) A part-solid nodule (synonym, semisolid nodule) consists of both ground-glass and solid soft-tissue attenuation components. (See also mass.)

nonspecific interstitial pneumonia, or NSIP

Pathology.—NSIP is characterized by a histologic pattern of uniform interstitial involvement by varying degrees of chronic inflammation or fibrosis. NSIP may be idiopathic or seen in other settings, including collagen vascular disease, hypersensitivity pneumonitis, drug-induced lung disease, infection, and immunodeficiency (including human immunodeficiency virus infection) (5).

CT scans.—NSIP has variable thin-
section CT appearances: The most frequent is ground-glass opacities with reticulation, traction bronchiectasis or bronchiolectasis, and little or no honeycombing (Fig 43). The distribution is usually basal and subpleural (95).

oligemia
Pathophysiology.—Oligemia is a reduction in pulmonary blood volume. Most frequently, this reduction is regional, but occasionally it is generalized. Regional oligemia is usually associated with reduced blood flow in the oligemic area.

Radiographs and CT scans.—Oligemia appears as a regional or widespread decrease in the size and number of identifiable pulmonary vessels (Fig 44), which is indicative of less than normal blood flow. (See also mosaic attenuation pattern, pulmonary blood flow redistribution.)

opacity
Radiographs and CT scans.—Opacity refers to any area that preferentially attenuates the x-ray beam and therefore appears more opaque than the surrounding area. It is a nonspecific term that does not indicate the size or pathologic nature of the abnormality. (See also parenchymal opacification, ground-glass opacity.)

organizing pneumonia
Pathology.—Organizing pneumonia manifests as a histologic pattern characterized by loose plugs of connective tissue in the airspaces and distal airways. Interstitial inflammation and fibrosis are minimal or absent. Cryptogenic organizing pneumonia, or COP, is a distinctive clinical disorder among the idiopathic interstitial pneumonias (5), but the histologic pattern of organizing pneumonia is encountered in many different situations, including pulmonary infection, hypersensitivity pneumonitis, and collagen vascular diseases.

Radiographs and CT scans.—Airspace consolidation is the cardinal feature of organizing pneumonia on chest radiographs and CT scans. In COP, the distribution is typically subpleural and basal (Fig 45) and sometimes bronchocentric (96). Other manifestations of organizing pneumonia include ground-glass opacity, tree-in-bud pattern, and nodular opacities (37).

panacinar emphysema
Pathology.—Panacinar emphysema involves all portions of the acinus and secondary pulmonary lobule more or less uniformly (42). It predominates in the lower lobes and is the form of emphysema associated with a1-antitrypsin deficiency.

CT scans.—Panacinar emphysema may coexist and merge with severe centrilobular emphysema. The appearance of featureless decreased attenuation may be indistinguishable from severe constrictive obliterative bronchiolitis (99). The term panlobular emphysema is synonymous. (See also emphysema.)
Paraseptal emphysema

Pathology.—Paraseptal emphysema is characterized by predominant involvement of the distal alveoli and their ducts and sacs. It is characteristically bounded by any pleural surface and the interlobular septa (42,43).

CT scans.—This emphysema is characterized by subpleural and peribronchovascular regions of low attenuation separated by intact interlobular septa (Fig 47), sometimes associated with bullae. The term distal acinar emphysema is synonymous. (See also emphysema.)

Parenchyma

Anatomy.—Parenchyma refers to the gas-exchanging part of the lung, consisting of the alveoli and their capillaries.

Radiographs and CT scans.—The portion of the lung exclusive of visible pulmonary vessels and airways.

Parenchymal band

Radiographs and CT scans.—A parenchymal band is a linear opacity, usually 1–3 mm thick and up to 5 cm long that usually extends to the visceral pleura (which is often thickened and may be retracted at the site of contact) (Fig 48). It reflects pleuroparenchymal fibrosis and is usually associated with distortion of the lung architecture. Parenchymal bands are most frequently encountered in individuals who have been exposed to asbestos (100,101).

Parenchymal opacification

Radiographs and CT scans.—Parenchymal opacification of the lungs may or may not obscure the margins of vessels and airway walls (45). Consolidation indicates that definition of these margins (excepting air bronchograms) is lost within the dense opacification, whereas ground-glass opacity indicates a smaller increase in attenuation, in which the definition of underlying structures is preserved (59). The more specific terms consolidation and ground-glass opacity are preferred. (See also consolidation, ground-glass opacity.)

Peribronchovascular interstitium

Anatomy.—The peribronchovascular interstitium is a connective-tissue sheath that encloses the bronchi, pulmonary arteries, and lymphatic vessels. It extends from the hila to the lung periphery.

Perilobular distribution

Anatomy.—The perilobular region comprises the structures bordering the periphery of the secondary pulmonary lobule.

CT scans.—This pattern is characterized by distribution along the structures that border the pulmonary lobules (ie, interlobular septa, visceral pleura, and vessels) (102). The term is most frequently used in the context of diseases (eg, perilobular organizing pneumonia) that are distributed mainly around the inner surface of the secondary pulmonary lobule (103) (Fig 49). This may resemble indistinct thickening of the interlobular septa.
perilymphatic distribution

**Anatomy.**—This pattern is characterized by distribution along or adjacent to the lymphatic vessels in the lung. The routes of lymphatics are found along bronchovascular bundles, in the interlobular septa, around larger pulmonary veins, and in the pleura; alveoli do not have lymphatics.

**CT scans.**—Abnormalities along the pathway of the pulmonary lymphatics—that is, in the perihilar, peribronchovascular, and centrilobular interstitium, as well as in the interlobular septa and subpleural locations—have a perilymphatic distribution (104). A perilymphatic distribution typically seen in sarcoidosis (Fig 50) and lymphangitic spread of cancer.

pleural plaque

**Pathology.**—A pleural plaque is a fibrohyaline, relatively acellular lesion arising predominantly on the parietal pleural surface, particularly on the diaphragm and underneath ribs (105). Pleural plaques are almost invariably the consequence of previous (at least 15 years earlier) asbestos exposure.

**Radiographs and CT scans.**—Pleural plaques are well-demarcated areas of pleural thickening, seen as elevated flat or nodular lesions that often contain calcification (Fig 51). Plaques are of variable thickness, range from less than 1 to approximately 5 cm in diameter, and are more easily identified on CT scans than on chest radiographs (106). An en face plaque may simulate a pulmonary nodule on chest radiographs. (See also pseudoplaque.)

platelike atelectasis

*See linear atelectasis.*

pneumatocele

**Pathology.**—A pneumatocele is a thin-walled, gas-filled space in the lung. It is most frequently caused by acute pneumonia, trauma, or aspiration of hydrocarbon fluid and is usually transient. The mechanism is believed to be a combination of parenchymal necrosis and check-valve airway obstruction (107).

**Radiographs and CT scans.**—A pneumatocele appears as an approximately round, thin-walled airspace in the lung (Fig 52).

pneumomediastinum

**Pathology.**—Pneumomediastinum is the presence of gas in mediastinal tissue outside the esophagus and tracheobronchial tree. It may be caused by
spontaneous alveolar rupture, with subsequent tracking of air along the bronchovascular interstitium into the mediastinum. Pneumomediastinum is particularly associated with a history of asthma, severe coughing, or assisted ventilation.

Radiographs and CT scans.—Pneumomediastinum appears as lucent streaks on chest radiographs, mostly vertically oriented (Fig 53). Some of these streaks may outline vessels and main bronchi. (See also pneumopericardium.)

pneumonia
Pathology.—Pneumonia is inflammation of the airspaces and/or interstitium (eg, due to infection, as in bacterial pneumonia). Infective pneumonia is characterized by exudate resulting in consolidation. The term is also used to refer to a number of noninfectious disorders of the lung parenchyma characterized by varying degrees of inflammation and fibrosis (eg, idiopathic interstitial pneumonias) (5).

pneumopericardium
Pathology.—Pneumopericardium is the presence of gas in the pericardial space. It usually has an iatrogenic, often surgical, origin in adults.

Radiographs and CT scans.—Pneumopericardium is usually distinguishable from pneumomediastinum because the lucency (low attenuation) caused by air does not extend outside the pericardial sac (Fig 54). (See also pneumomediastinum.)

cardiac tamponade
Pathophysiology.—Cardiac tamponade is the accumulation of fluid in the pericardial space, with resultant compression of the cardiac chambers. Symptoms include dyspnea, hypotension, and muffled heart sounds. On chest radiography, the heart may appear enlarged or displaced. Echocardiography is the diagnostic test of choice.

pneumothorax and tension pneumothorax
Pathophysiology.—Pneumothorax refers to the presence of gas in the pleural space. Qualifiers include spontaneous, traumatic, diagnostic, and tension. Tension pneumothorax is the accumulation of intrapleural gas under pressure. In this situation, the ipsilateral lung will, if normal, collapse completely; however, a less than normally compliant lung may remain partially inflated.

Radiographs and CT scans.—On chest radiographs, a visceral pleural edge is visible (Fig 55) unless the pneumothorax is very small or the pleural edge is not tangential to the x-ray beam. Tension pneumothorax may be associated with considerable shift of the mediastinum and/or depression of the hemidiaphragm. Some shift can occur without tension because the pleural pressure in the presence of pneumothorax becomes atmospheric, while the pleural pressure in the contralateral hemithorax remains negative.

progressive massive fibrosis
Pathology.—This condition is caused by slow-growing conglomeration of dust particles and collagen deposition in individuals (mostly coal workers) heavily exposed to inorganic dust (108).

Radiographs and CT scans.—Progressive massive fibrosis manifests as masslike lesions, usually bilateral and in the upper lobes (Fig 56). Background nodular opacities reflect accompanying pneumoconiosis, with or without emphysematous destruction adjacent to the massive fibrosis (109). Lesions similar to progressive massive fibrosis sometimes occur in other conditions, such as sarcoidosis and talcosis (109,110).
pseudocavity

CT scans.—A pseudocavity appears as an oval or round area of low attenuation in lung nodules, masses, or areas of consolidation that represent spared parenchyma, normal or ectatic bronchi, or focal emphysema rather than cavitation. These pseudocavities usually measure less than 1 cm in diameter. They have been described in patients with adenocarcinoma (Fig 57), bronchioloalveolar carcinoma (111), and benign conditions such as infectious pneumonia.

pseudoplaque

CT scans.—A pseudoplaque is a pulmonary opacity contiguous with the visceral pleura formed by coalescent small nodules. It simulates the appearance of a pleural plaque. This entity is encountered most commonly in sarcoidosis (Fig 58), silicosis, and coal-worker’s pneumoconiosis (86).

pulmonary blood flow redistribution

Pathophysiology.—Pulmonary blood flow redistribution refers to any departure from the normal distribution of blood flow in the lungs that is caused by an increase in pulmonary vascular resistance elsewhere in the pulmonary vascular bed.

Radiographs and CT scans.—Pulmonary blood flow redistribution is indicated by a decrease in the size and/or number of visible pulmonary vessels in one or more lung regions (Fig 59), with a corresponding increase in number and/or size of pulmonary vessels in other parts of the lung. Upper lobe blood diversion in patients with mitral valve disease is the archetypal example of redistribution (112,113).

respiratory bronchiolitis–interstitial lung disease, or RB-ILD

Pathology.—RB-ILD is a smoking-related disease characterized by inflammation (predominantly by macrophages) of the respiratory bronchioles and peribronchiolar alveoli (5), sometimes with elements of or overlap with nonspecific and desquamative interstitial pneumonias (114).

CT scans.—RB-ILD typically manifests as extensive centrilobular micronodules and patchy ground-glass opacity typical of respiratory bronchiolitis–interstitial lung disease.

Figure 57: Transverse CT scan shows pseudocavitation in an adenocarcinoma.

Figure 58: Transverse CT scan shows pseudoplaques (arrows) in a patient with sarcoidosis.

Figure 59: Chest radiograph shows redistribution of blood flow to upper lung zones.

Figure 60: Transverse CT scan shows centrilobular micronodules and patchy ground-glass opacity typical of respiratory bronchiolitis–interstitial lung disease.
**reticular pattern**

*Radiographs and CT scans.*—On chest radiographs, a reticular pattern is a collection of innumerable small linear opacities that, by summation, produce an appearance resembling a net (synonym: reticulation) (Fig 61). This finding usually represents interstitial lung disease. The constituents of a reticular pattern are more clearly seen at thin-section CT, whether they are interlobular septal thickening, intralobular lines, or the cyst walls of honeycombing. (*Reticular pattern and honeycombing should not be considered synonymous. See also honeycombing.*)

**reticulonodular pattern**

*Radiographs and CT scans.*—A combined reticular and nodular pattern, the reticulonodular pattern is usually the result of the summation of points of intersection of innumerable lines, creating the effect on chest radiographs of superimposed micronodules (Fig 62). The dimension of the nodules depends on the size and number of linear or curvilinear elements. (*See also reticular pattern.*) On CT scans, the pattern appears as a concurrence of reticular and micronodular patterns. The micronodules can be situated in the center of the reticular elements (eg, centrilobular micronodules) or superimposed on the linear opacities (eg, septal micronodules).

**reversed halo sign**

*CT scans.*—The reversed halo sign is a focal rounded area of ground-glass opacity surrounded by a more or less complete ring of consolidation (Fig 63). A rare sign, it was initially reported to be specific for cryptogenic organizing pneumonia (117,118) but was subsequently described in patients with paracoccidioidomycosis (119). Similar to the halo sign, this sign will probably lose its specificity as it is recognized in other conditions. (*See also halo sign.*)

**right paratracheal stripe**

*Anatomy and radiographs.*—The right paratracheal stripe is a vertical, linear, soft-tissue opacity less than 4 mm wide. It corresponds to the right tracheal wall, contiguous mediastinal tissues, and adjacent pleura (Fig 64). The stripe is 3–4 cm in height and extends from approximately the level of the medial ends of the clavicles to the right tracheobronchial angle on a frontal chest radiograph (120). It is seen in up to 94% of adults but is widened or absent in individuals with abundant mediastinal fat. The commonest pathologic cause of widening, deformity, or obliteration of this stripe is paratracheal lymph node enlargement.
rounded atelectasis

**Pathology.**—Rounded atelectasis is rounded collapsed lung associated with invaginated fibrotic pleura and thickened and fibrotic interlobular septa. Most frequently, it is the consequence of an asbestos-induced exudative pleural effusion with resultant pleural scarring (121), but it may occur with any cause of pleural fibrosis.

**Radiographs and CT scans.**—Individual segments cannot be precisely delineated on chest radiographs and CT scans, and their identification is made inferentially on the basis of the position of the supplying segmental bronchus and artery. When occasionally present, intersegmental fissures help to identify segments.

**septal line**
See interlobular septum.

**septal thickening**
See interlobular septal thickening.

**segment**

**Anatomy.**—A segment is the unit of a lobe ventilated by a segmental bronchus, perfused by a segmental pulmonary artery, and drained by an intersegmental pulmonary vein. There are two to five segments per lobe.

**Radiographs and CT scans.**—Individual segments cannot be precisely delineated on chest radiographs and CT scans, and their identification is made inferentially on the basis of the position of the supplying segmental bronchus and artery. When occasionally present, intersegmental fissures help to identify segments.

**silhouette sign**

**Radiographs.**—The silhouette sign is the absence of depiction of an anatomic soft-tissue border. It is caused by consolidation and/or atelectasis of the adjacent lung (Fig 67), by a large mass, or by contiguous pleural fluid (128,129). The silhouette sign results from the juxtaposition of structures of similar radiographic attenuation. The sign actually refers to the absence of a silhouette. It is not always indicative of disease (eg, unexplained absence of right border of heart is seen in association with pectus excavatum and sometimes in healthy individuals).

**small-airways disease**

**Pathology.**—This phrase is an arbitrary term used more frequently in thin-section CT descriptions than in the pathophysiology literature, where it was first coined (130). *Small-airways disease* now generally refers to any condition affecting the bronchioles, whereas...
bronchiolitis more specifically describes inflammation of the bronchioles (131).

CT scans.—Small airways are considered to be those with an internal diameter smaller than or equal to 2 mm and a normal wall thickness of less than 0.5 mm (32). Small-airways disease is manifest on CT scans as one or more of the following patterns: mosaic attenuation, air trapping, centrilobular micronodules, tree-in-bud pattern, or bronchiolectasis.

subpleural curvilinear line

CT scans.—This finding is a thin curvilinear opacity, 1–3 mm in thickness, lying less than 1 cm from and parallel to the pleural surface (Fig 68). It corresponds to atelectasis of normal lung if seen in the dependent posteroinferior portion of lung of a patient in the supine position and is subsequently shown to disappear on CT sections acquired with the patient prone. It may also be encountered in patients with pulmonary edema (132) or fibrosis (other signs are usually present). Though described in the context of asbestosis, this finding is not specific for asbestosis.

traction bronchiectasis and traction bronchiolectasis

CT scans.—Traction bronchiectasis and traction bronchiolectasis respectively represent irregular bronchial and bronchiolar dilatation caused by surrounding retractile pulmonary fibrosis (130). Dilated airways are usually identifiable as such (Fig 69) but may be seen as cysts (bronchi) or microcysts (bronchioles in the lung periphery). The juxtaposition of numerous cystic airways may make the distinction from “pure” fibrotic honeycombing difficult.

tree-in-bud pattern

CT scans.—The tree-in-bud pattern represents centrilobular branching structures that resemble a budding tree. The pattern reflects a spectrum of endo- and peribronchiolar disorders, including mucoid impaction, inflammation, and/or fibrosis (134,135) (Fig 70). This pattern is most pronounced in the lung periphery and is usually associated with abnormalities of the larger airways. It is particularly common in diffuse panbronchiolitis (136), endobronchial spread of mycobacterial infection (137), and cystic fibrosis. A similar pattern is a rare manifestation of arteriolar (microangiopathic) disease (138).
usual interstitial pneumonia, or UIP

Pathology.—UIP is a histologic pattern of pulmonary fibrosis characterized by temporal and spatial heterogeneity, with established fibrosis and honeycombing interspersed among normal lung. Fibroblastic foci with fibrotic destruction of lung architecture, often with honeycombing, are the key findings (5). The fibrosis is initially concentrated in the lung periphery. UIP is the pattern seen in idiopathic pulmonary fibrosis, but can be encountered in diseases of known cause (eg, some cases of chronic hypersensitivity pneumonitis).

Radiographs and CT scans.—Honeycombing with a basal and subpleural distribution (Fig 71) is regarded as pathognomonic (63,65), but not all cases of biopsy-proved UIP have this distinctive CT pattern.

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