The role of high resolution computed tomography in the diagnosis of interstitial lung disease

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Until recently the chest radiograph has been the only imaging technique used in the assessment of patients with suspected diffuse lung disease. In this context the chest radiograph is less than ideal. Problems arise with both false negative and false positive results. It is well established that the chest radiograph may appear entirely normal in up to 10% of patients with biopsy proved diffuse lung disease of various causes,1 and a poor quality chest radiograph, especially of an obese patient, may misleadingly raise the spectre of diffuse lung disease. The two dimensional nature of a chest radiograph dictates that there is superimposition of structures over the lungs and it has been estimated that up to 40% of the lungs is obscured in normal subjects. This obscuration will be further compounded by the presence of pleural thickening or an effusion. Even when the chest radiograph shows definite evidence of diffuse lung disease McCloud has pointed out that “the chest radiograph is often non-specific. Various radiographic patterns, together with their predominant location, correlate statistically with their pathologic entities but in individual cases the chest roentgenogram is rarely diagnostic.”2

Until recently the main use for computed tomography has been detection of pulmonary nodules and, despite some pioneering work by Kree1 in the early 1980s, it has had a limited role in the investigation of diffuse lung disease. Improvements in computed tomography scanner technology, notably in terms of spatial resolution and shorter scan times, have led to renewed interest in the application of computed tomography to show the fine morphological detail of the lung. Research in this area is particularly active and is reflected in the large volume of descriptive work on the computed tomographic appearances of many diffuse lung diseases that has appeared over the last five years; there are no signs that this period of research is over. The technique of high resolution computed tomography, used to show the lung parenchyma,3 is sufficiently different from conventional computed tomography to warrant a brief description.

Conventional computed tomography of the thorax uses contiguous 1 cm sections; in this way the entire lungs are included in one study. This protocol is widely used when a comprehensive examination of the lungs is required—for example, in the search for metastases. The volume averaging that occurs within the 1 cm thickness of the scan, however, substantially reduces the ability of conventional computed tomography to resolve small structures. For high resolution computed tomography the section thickness is reduced to 1-3 mm and a different software reconstruction of the image is used to improve spatial resolution (figs 1a and 1b). These scans are interspaced by at least 1 cm. Where 3 mm sections are taken 1 cm apart the radiation dose to the breast is reduced to about half that of conventional computed tomography. None the less, the radiation burden inherent in high resolution computed tomography is considerable and in a complete study using 3 mm sections every 1 cm the dose is roughly 40 times that received from a single chest radiograph.4

Because high resolution computed tomography includes only very short segments of pulmonary vessels these narrow sections may be misinterpreted as showing a nodular pattern. For this reason some radiologists advocate conventional computed tomography before high resolution computed tomography but this obviously adds to the radiation dose. Because, by definition, diffuse lung disease is widespread many radiologists now perform a limited number of high resolution computed tomography scans, perhaps as few as six, in such patients. In this way not only is the radiation dose restricted but the cost of the examination is not much more than that of a posteroanterior and lateral chest radiograph. It has been estimated in the United States that high resolution computed tomography used in this way costs $180, compared with $90 for a posteroanterior and lateral chest radiograph. In the same comparison a transbronchial biopsy with bronchoalveolar lavage costs $1500 and an open lung biopsy $5000 (R A Webb, personal communication). Computed tomography is becoming more widely available throughout the United Kingdom and most computed tomography machines installed over the last three years are able to provide high resolution images.

The spatial resolution of the latest gener-
Figure 1  
(a) Conventional 10 mm width computed tomographic section through the upper lobes (normal). (b) High resolution 3 mm width computed tomographic section at the same level. Note the different appearance of the blood vessels and bronchi.

Attempted discrimination of differences on chest radiographs. Conventional chest radiography is limited in its ability to differentiate between airspace and interstitial disease. It often fails to distinguish whether thickening of the interstitium and bronchovascular structures are evenly confluent, or if there is a predominantly interstitial pattern of disease. The most common abnormality on the chest radiograph is interstitial thickening, with the additional pattern of abnormal shadowing due to airspace disease.

Compared to conventional tomography, high resolution computed tomography provides important information about the interstitial and airspace components of the lung. This technique is able to resolve structures as small as 200 μm, allowing visualization of the secondary pulmonary lobule and its associated septa. The interlobular septum measures 100 μm across in normal individuals and is therefore more often identified in pathological states such as thickening of the interstitium. The secondary pulmonary lobule itself is an irregular polyhedron and contains a connective tissue sheath that surrounds the bronchovascular bundle. This core may be visible as a small dot in the centre of the lobule.

High resolution computed tomography through the upper lobes shows unilateral thickening of the bronchovascular bundles and interlobular septa, features from the pattern on a chest radiograph have largely been discredited. This is partly because the abnormal shadowing on a radiograph results from summation of numerous foci; the chest radiograph has been fancifully described as "the two-dimensional projectional sum of several thousand histologic slices." Another compelling reason is the overwhelming frequency with which interstitial and airspace lesions coexist in any given disease (see fig 2). Nevertheless, high resolution computed tomography does show interstitial thickening precisely in the few diseases in which the pathological process is largely confined to the interstitium. An example is shown in figure 4; the chest radiograph of this patient showed non-specific shadowing in the right upper zone. Despite extensive investigation a diagnosis was not reached. High resolution computed tomography through the upper lobes shows unilateral thickening of the bronchovascular bundles and interlobular septa, features
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Figure 2. (a) Polygonal outlines representing thickened interlobular septa (arrowheads) in silicosis. The background nodularity is more obvious on sections through the upper zones. (b) Thickening of the interlobular septa in areas of airspace consolidation due to alveolar proteinosis.

characteristic of lymphangitis carcinomatosa.\textsuperscript{11} From the pathological descriptions of lymphangitis the high resolution computed tomography appearances can be predicted and the same applies to pulmonary sarcoidosis.\textsuperscript{1} There are subtle but definite features that differentiate the high resolution computed tomography appearances of sarcoidosis from lymphangitis carcinomatosa: in sarcoidosis the nodular thickening of the bronchovascular bundles is more pronounced and thickening of the interlobular septa less obvious (fig 5). Nevertheless, as with chest radiography, pulmonary sarcoidosis has many guises and a wide range of high resolution computed tomography appearances has been reported\textsuperscript{11} (fig 6). Fibrosing alveolitis, whether “lone” or associated with other diseases, has been shown to have a characteristic computed tomographic appearance. The crescentic subpleural distribution concentrated in the posterobasal segments of the lower lobes is virtually pathognomonic\textsuperscript{14,15} (fig 7). Not surprisingly, given this basal distribution, high resolution computed tomography will show disease despite a normal chest radiograph.\textsuperscript{16} Scans taken in the prone position may be helpful in distinguishing parenchymal opacification due to increased perfusion of the dependent parts of the lung from minimal fibrosing alveolitis. As the fibrosis progresses a widespread, coarse reticular pattern supervenes, resulting in a non-specific appearance of end stage pulmonary fibrosis. In the clinical context of unexplained deterioration of a patient with fibrosing alveolitis high resolution computed tomography may be of value as it may reveal other disease—for example, a bronchogenic carcinoma. High resolution com-
Figure 3 Intralobular bronchovascular bundles visible as minute central dots (arrows) in this case of subacute extrinsic allergic alveolitis.

Figure 4 Lymphangitis carcinomatosa: in the right lung there is thickening of the bronchovascular bundles and prominent subpleural interlobular septa. There is a pleural plaque posteriorly in the left chest.

Computed tomography may also show coexisting emphysema, which may be impossible to detect by chest radiography. There is increasing evidence of a high degree of correlation between high resolution computed tomography appearances and the amount of disease activity in fibrosing alveolitis in terms of the proportion of cellular infiltrate to established fibrosis found histologically. The observation that amorphous parenchymal opacification represents increased cellularity, in both the airspaces and the interstitium (fig 8), whereas the reticular pattern is the result of fibrosis (fig 7) appears, with some provisos, to be true. No other diagnostic technique (including lung function tests) short of biopsy allows this
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Figure 5  Sarcoidosis, with nodules predominantly concentrated around bronchovascular bundles.

Figure 6  Sarcoidosis: a less common pattern with irregular areas of confluent parenchymal opacification (prone scan).

distinction to be made reliably.

The superior sensitivity of high resolution computed tomography for detecting asbestosis is now undisputed, particularly in those patients with extensive pleural disease, which obscures the underlying lung.21 22 High resolution computed tomography, however, has opened a window on minor degrees of pleuroparenchymal abnormality whose long term significance is as yet unknown. It will be some years before the natural history of these changes is established. Other conditions in which high resolution computed tomography frequently shows parenchymal abnormality in
the face of a normal chest radiograph include subacute extrinsic allergic alveolitis23 (fig 9) and drug induced lung damage.24 The high resolution computed tomography patterns of most of the rarer diffuse lung diseases have now been described in detail. These include the pulmonary manifestations of lymphangioleiomyomatosis,25,26 histiocytosis X,27 and tuberous sclerosis.28 Although these patterns may not in themselves be unerringly diagnostic, in a given clinical context they will often confirm the diagnosis and at the same time exclude other suspected diseases, such as emphysema. The example of lymphangioleiomyomatosis shown in figure 10 demonstrates numerous cystic air spaces of uniform...
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Figure 9 Subacute extrinsic allergic alveolitis: global parenchymal opacification due to infiltration of the interstitium and airspaces. A concurrent chest radiograph was normal.

Figure 10 Lymphangioleiomyomatosis: well defined cystic airspaces. Note the preservation of a normal vascular pattern. The chest radiograph showed minimal and non-specific changes.

size with fine but definable walls (in contrast to the areas of lung destruction seen in emphysema, which do not have discernible margins). This woman with lymphangioleiomyomatosis had a spontaneous pneumothorax during pregnancy; the underlying lungs appeared questionably abnormal on the chest radiograph. In this instance a definite diagnosis was made from the high resolution computed tomography appearances without resorting to an open lung biopsy.

Although the sensitivity of high resolution computed tomography in detecting minor degrees of diffuse lung disease is not seriously challenged by any other imaging technique currently available, its precise role in the dyspnoic patient with a normal chest radiograph and abnormal results in lung function tests remains to be determined. It seems likely that high resolution computed tomography will obviate the need for lung biopsy by providing a specific diagnosis in some of these patients. In patients needing a biopsy high resolution computed tomography should be used to provide guidance to the site most likely to yield satisfactory histological material.

The utility of high resolution computed tomography compared with chest radiography
in predicting a specific diagnosis of diffuse lung disease has been elegantly demonstrated in a study from Vancouver.29 The radiographs and computed tomograms of 118 patients, with widely varying diagnoses proved by biopsy, were assessed by three observers without knowledge of any clinical data, other than the age of the patient. The observers listed the three most likely diagnoses in order of probability and recorded their level of confidence for the first diagnosis. Overall, regardless of confidence levels, the correct diagnosis was made in 57% of cases with chest radiography and 76% of cases with high resolution computed tomography. High confidence was achieved in 23% of chest radiograph readings and 49% of high resolution computed tomography readings; of these readings the correct diagnosis were made in 77% and 93% respectively. The implications are that high resolution computed tomography will allow a confident and specific diagnosis of diffuse lung disease to be made considerably more often and more accurately than with chest radiography.

As an aside to the main purpose of this study, Mathieson et al showed that HRCT correctly predicted whether transbronchial or open lung biopsy was more appropriate in 87% of cases, compared with 65% when chest radiography was used (the criterion being that if sarcoidosis or lymphangitis carcinomatosa was the favoured diagnosis then bronchial biopsy was indicated). This is an important study, which should be repeated in the light of increasing experience with high resolution computed tomography.

Now that the correlations between the high resolution computed tomography appearances and the pathological features of most diffuse lung diseases have been thoroughly studied, other areas receiving attention are functional correlations between the extent of disease as shown by high resolution computed tomography and the results of lung function tests. Longitudinal studies of the rate of change by diffuse lung disease detected by high resolution computed tomography compared with that shown by other monitoring techniques, both imaging and functional, are also needed.

The availability and sophistication of computed tomography machines capable of producing high resolution images continues to increase, but skill in interpreting these images is not increasing at the same rate and remains limited to a few centres. Given the obvious utility of high resolution computed tomography in diffuse lung disease (summarised in the table), chest physicians and radiologists should now consider high resolution computed tomography a routine and indispensable part of the investigation and assessment of a patient with suspected diffuse lung disease.

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