

# Chapter 33

## Interstitial Pneumonia



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Describe the classic CXR appearance of acute interstitial pneumonia [3].	<i>Diffuse heterogeneous opacities.</i>
Describe the classic CT appearance of acute interstitial pneumonia [3].	<i>Extensive symmetric ground-glass opacities associated with traction bronchiectasis.</i>
Which imaging modality is preferred for characterizing diffuse pulmonary disease? [3]	<i>CT.</i>
What is a key radiologic difference between acute interstitial pneumonia and desquamative interstitial pneumonia? [4]	<i>DIP does not have traction bronchiectasis.</i>
Bilateral, lower zone-predominant ground-glass opacities ± subpleural intralobular lines are suggestive of which type of interstitial pneumonia? [5]	<i>Desquamative interstitial pneumonia.</i>

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What are the common demographics for desquamative interstitial pneumonia? [5]	<i>Smokers; M:F, 2:1; age, 40–60.</i>
Bilateral, diffuse, ground-glass opacities in an immunocompromised patient suggests which pathogen? [1–3]	<i>Pneumocystis jirovecii.</i>
What is the best imaging modality for pneumocystis pneumonia? [6]	<i>HRCT.</i>
What is the best diagnostic clue on CT for lymphoid interstitial pneumonia?	<i>Ground-glass opacities ± pulmonary cysts.</i>
What is seen on CXR of lymphoid interstitial pneumonia?	<i>Basilar reticular/reticulonodular opacities.</i>

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## References

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