

CLINICAL IMAGE

Hepatopulmonary syndrome complicated by interstitial pneumonia and obesity with normal contrast echocardiography

Akihito Okazaki^{1,2}  | Kensuke Fujioka³¹Department of Respiratory Medicine, Koseiren Takaoka Hospital, Takaoka, Japan²Department of Respiratory Medicine, Kaga Medical Center, Kaga, Japan³Department of Cardiovascular Medicine, Koseiren Takaoka Hospital, Takaoka, Japan**Correspondence**

Akihito Okazaki, Department of Respiratory Medicine, Kaga Medical Center, Ri-36, Sakumi-machi, 922-8522 Kaga, Ishikawa, Japan.
Email: akihitookazaki1017@gmail.com

Funding information

This research was not supported by any specific grant from any funding agency in the public, commercial, or non-profit sectors. Therefore, no funders were involved in the design of the study, the collection, analysis, and interpretation of the data writing of the manuscript, or the decision to submit the manuscript for publication

Abstract

Diagnosis of hepatopulmonary syndrome complicated by interstitial pneumonia and obesity is difficult because these complications can cause hypoxia. Such patients may not present with typical contrast echocardiography findings.

KEYWORDS

hepatopulmonary syndrome, interstitial pneumonia, perfusion lung scintigraphy

1 | CASE PRESENTATION

Hepatopulmonary syndrome (HPS) presents with three features: liver disease, intrapulmonary blood vessel dilation, and hypoxemia. HPS complicated by interstitial pneumonia and obesity is difficult to diagnose because these complications can independently cause hypoxia. Multiple modalities, including pulmonary function test, contrast-enhanced echocardiography, and perfusion scintigraphy, should be combined for accurate diagnosis.

A 60-year-old man with alcoholic liver cirrhosis and obesity (BMI, 27.3) presented with exertional dyspnea. Chest radiography revealed bilateral diffuse interstitial opacities (Figure 1). Laboratory examination and pulmonary function tests revealed arterial hypoxemia (PaO₂, 53.0 mmHg),

elevated KL-6 (2,087 U/mL), and reduced % DLCO (30.8%). Total lung capacity, residual volume, and forced vital capacity were within the normal range. Echocardiography showed no intracardiac shunt, and the microbubble test results were negative. Perfusion lung scanning revealed no evidence of pulmonary embolism, but a severe right-to-left shunt was identified (Figure 1). Therefore, the patient was diagnosed with hepatopulmonary syndrome (HPS).

2 | DISCUSSION AND CONCLUSION

This case provides two important clinical suggestions. First, diagnosis of HPS complicated by interstitial

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

© 2021 The Authors. *Clinical Case Reports* published by John Wiley & Sons Ltd.

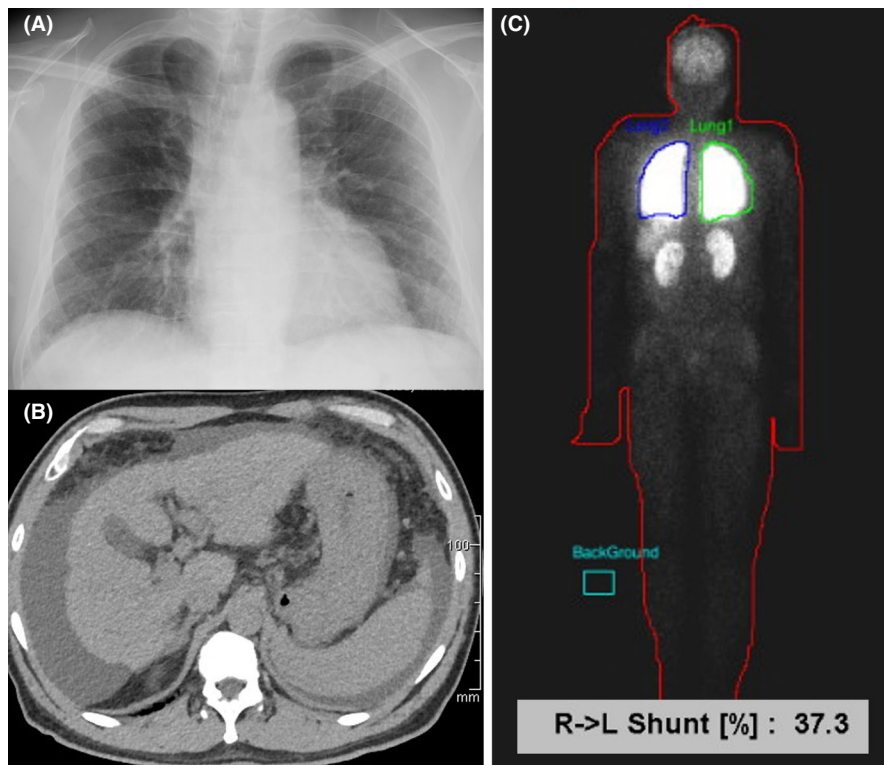


FIGURE 1 Chest radiography revealing bilateral diffuse interstitial opacities (A). Total lung capacity, residual volume, and forced vital capacity are within normal range, and % DLCO is decreased to 30.8%. Computed tomography revealing liver atrophy, surface irregularities, splenomegaly, and ascites (B). Lung perfusion scintigraphy with ^{99m}Tc -MAA revealing accumulation in the brain, kidneys, and gastrointestinal tract (shunt=37.3%) (C). Defects in ^{99m}Tc -MAA are absent in the lungs. Home oxygen therapy is initiated, and liver transplantation is currently under consideration

pneumonia and obesity is difficult. Diffusion impairment due to pulmonary fibrosis and obesity-induced thoracic compression can also cause hypoxia. The presence of decreased DLCO and absence of abnormalities in lung volume fractions lead to suggest the possibility of HPS.

Second, normal contrast-enhanced echocardiography (CEE) findings may be obtained despite the presence of a high degree shunt. Some reports indicated that perfusion lung scanning was more sensitive than CEE.^{1,2} Multiple modalities, including pulmonary perfusion scintigraphy, should be combined for accurate diagnosis.

ACKNOWLEDGEMENTS

None.

CONFLICTS OF INTEREST

None.

AUTHOR CONTRIBUTIONS

AO was responsible for conception, design, drafting, image modification, and finalizing the manuscript. KF was responsible for demonstrating contrast echocardiography. All authors read and approved the final manuscript.

CONSENT

Informed consent for the publication and related images was obtained from the patient.

DATA AVAILABILITY STATEMENT

No datasets were generated or analyzed during this case report.

ORCID

Akihito Okazaki  <https://orcid.org/0000-0002-5426-9035>

REFERENCES

1. El-Shabrawi MH, Omran S, Wageeh S, et al. (99m) technetium-macroaggregated albumin perfusion lung scan versus contrast enhanced echocardiography in the diagnosis of the hepatopulmonary syndrome in children with chronic liver disease. *Eur J Gastroenterol Hepatol.* 2010;22:1006-1012.
2. Alipour Z, Armin A, Mohamadi S, et al. Hepatopulmonary syndrome with right-to-left shunt in cirrhotic patients using macro-aggregated albumin lung perfusion scan: comparison with contrast echocardiography and association with clinical data. *Mol Imaging Radionucl Ther.* 2020;29:1-6.

How to cite this article: Okazaki A, Fujioka K. Hepatopulmonary syndrome complicated by interstitial pneumonia and obesity with normal contrast echocardiography. *Clin Case Rep.* 2021;9:e05064. <https://doi.org/10.1002/ccr3.5064>