

CLINICAL IMAGE

Pericostal tuberculosis in a patient with systemic sclerosis: The relationship of two rare diseases

Naoho Takizawa¹  | Tetsushi Mizutani² | Yoshiro Fujita¹¹Department of Rheumatology, Chubu Rosai Hospital, Nagoya, Japan²Department of Surgery, Chubu Rosai Hospital, Nagoya, Japan**Correspondence**

Naoho Takizawa, Department of Rheumatology, Chubu Rosai Hospital, 1-10-6 Komei, Minato-ku, Nagoya 455-8530, Japan.

Email: ttkkzww5959@gmail.com

Abstract

Regardless of immunosuppressant use, physicians should be aware of pulmonary and extra-pulmonary tuberculosis in patients with autoimmune disease including systemic sclerosis, especially if they follow unusual clinical courses.

KEY WORDS

Mycobacterium tuberculosis, scleroderma, systemic sclerosis, tuberculosis

1 | INTRODUCTION

Autoimmune diseases including systemic sclerosis (SSc) increase risk of developing TB. Pericostal tuberculosis (TB) is a rare presentation of skeletal TB. This case report describes pericostal TB in a SSc patient and emphasizes significance of suspecting pulmonary and extra-pulmonary TB when patients with autoimmune disease follow atypical clinical courses.

An 83-year-old woman presented with a left pericostal painful mass which she had had for a year. She was diagnosed with systemic sclerosis (SSc) by skin thickening and a positive result of anti-centromere antibody at age 75, however, she did not take any immunosuppressants. An abdominal CT showed a pericostal mass (Figure 1). We performed a needle biopsy, and the result was negative for Ziehl-Neelsen stain, but both the PCR and culture of tuberculosis (TB) from the drain were positive. She was diagnosed with pericostal TB, and we started a combination treatment with isoniazid, rifampicin, pyrazinamide, and ethambutol. Her chest pain resolved quickly and a repeat biopsy culture was done one month after initiation of treatment, and the result was negative. Pericostal TB is a rare presentation of skeletal TB and is thought to be caused by an extension of a TB infection of

the intercostal lymph nodes.¹ It has been reported that the risk of TB in SSc patients is 2.8 times higher than those in the general population.² The increased risk of developing TB in patients with autoimmune disorders may be due to an immune abnormality itself or immunosuppressants.² Regardless



FIGURE 1 Abdominal contrast-enhanced CT. White arrow showed pericostal mass with central hypoattenuation

This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

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

of immunosuppressant use, physicians should be aware of TB in SSc patients.

ACKNOWLEDGEMENTS

Published with written consent of the patient.

CONFLICT OF INTEREST

None declared.

AUTHOR CONTRIBUTIONS

Naoho Takizawa: wrote the initial draft, reviewed the literature, revised manuscript, and approved the final version. Tetsushi Mizutani: reviewed the literature and revised the manuscript. Yoshiro Fujita: reviewed the literature, revised the manuscript, and approved the final version.

ETHICAL STATEMENT

Written informed consent was obtained from the patient who participated in this study. This case report did not receive any funding. Authors have access to all source data for this case report.

DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available on request from the corresponding author. The data are not publicly available due to privacy or ethical restrictions.

ORCID

Naoho Takizawa  <https://orcid.org/0000-0003-3744-3537>

REFERENCES

1. Prasoon D. Tuberculosis of the intercostal lymph nodes. *Acta Cytol.* 2003;47:51-55.
2. Ou SM, Fan WC, Chou KT, et al. Systemic sclerosis and the risk of tuberculosis. *J Rheumatol.* 2014;41:1662-1669.

How to cite this article: Takizawa N, Mizutani T, Fujita Y. Pericostal tuberculosis in a patient with systemic sclerosis: The relationship of two rare diseases. *Clin Case Rep.* 2021;00:e04563. <https://doi.org/10.1002/ccr3.4563>