

**[ CASE REPORT ]**

# **Radiologically Undetectable Latent Pulmonary Lymphocyte Infiltration Suggesting the Need for Early Glucocorticoid Therapy: An Autopsy Case of Rapidly Progressive COVID-19 Pneumonia with Lymphopenia**

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## **Abstract:**

We herein report a fatal case of COVID-19 pneumonia with rapid progression of respiratory failure and lymphopenia. Excessive recruitment and sequestration of lymphocytes in the lung were suggested as the pathophysiology underlying COVID-19-associated lymphopenia. Interestingly, the autopsy in this case revealed lymphocytic infiltration in the lungs even at sites that appeared normal on autopsy imaging. These findings suggest that in COVID-19 cases with risk factors of severe exacerbation, early glucocorticoid administration should be considered, especially if lymphopenia is present, even if the imaging findings show only mild abnormalities.

**Key words:** COVID-19 pneumonia, autopsy, lymphocytic infiltration, lymphopenia

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## **Introduction**

Severe acute respiratory dysfunction often occurs in patients with coronavirus disease 2019 (COVID-19). Glucocorticoid therapy is indicated for the treatment of COVID-19 (1), but there is no clear consensus as to when to initiate it. We herein report a fatal case of COVID-19 with rapid progression of respiratory failure. An autopsy provided pathological and radiological findings supporting the efficacy of early glucocorticoid administration for COVID-19.

## **Case Report**

An 85-year-old man with chronic kidney disease and schizophrenia was transferred to our hospital with a suspected SARS-CoV-2 infection. He had been admitted to the previous hospital for schizophrenia during an outbreak of COVID-19 in his ward. On presentation, he had a two-day history of cough and a fever (day 1), and a positive reverse

transcription-polymerase chain reaction test confirmed COVID-19.

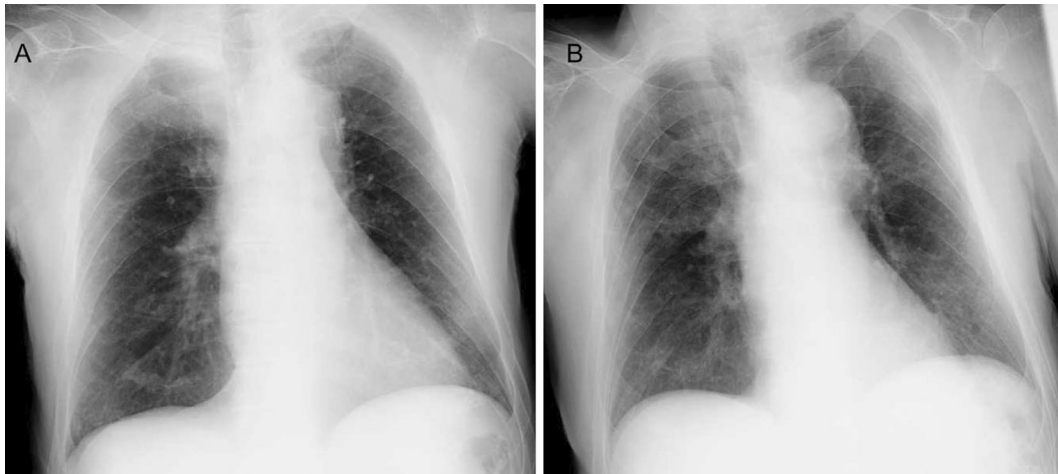
A physical examination revealed blood pressure 162/90 mmHg, heart rate 83/min, temperature 38.8°C, respiratory rate 20/min, and oxygen saturation 97% on room air. Laboratory tests at admission showed leukopenia but no increase in serum CRP levels. Chest X-ray showed slight, bilateral ground-glass lesions (Fig. 1A). The fever persisted after the transfer, and the serum CRP level increased. On hospitalization day 6, his SpO<sub>2</sub> decreased, and oxygen administration (1 L/min) was started. On hospitalization day 8, his oxygen demand increased to 2 L/min. Chest X-ray showed exacerbation of the ground-glass opacity (Fig. 1B). Dexamethasone was not used to treat COVID-19 due to the lack of any established recommendations at the time (2).

On the evening of hospitalization day 9, oxygen demand rapidly increased from 2 to 4 L/min. Given the grave condition of the patient, after careful discussion with the patient's family, the decision was made not to resuscitate. From 6 am on hospitalization day 10, tachypnea with a respiratory rate

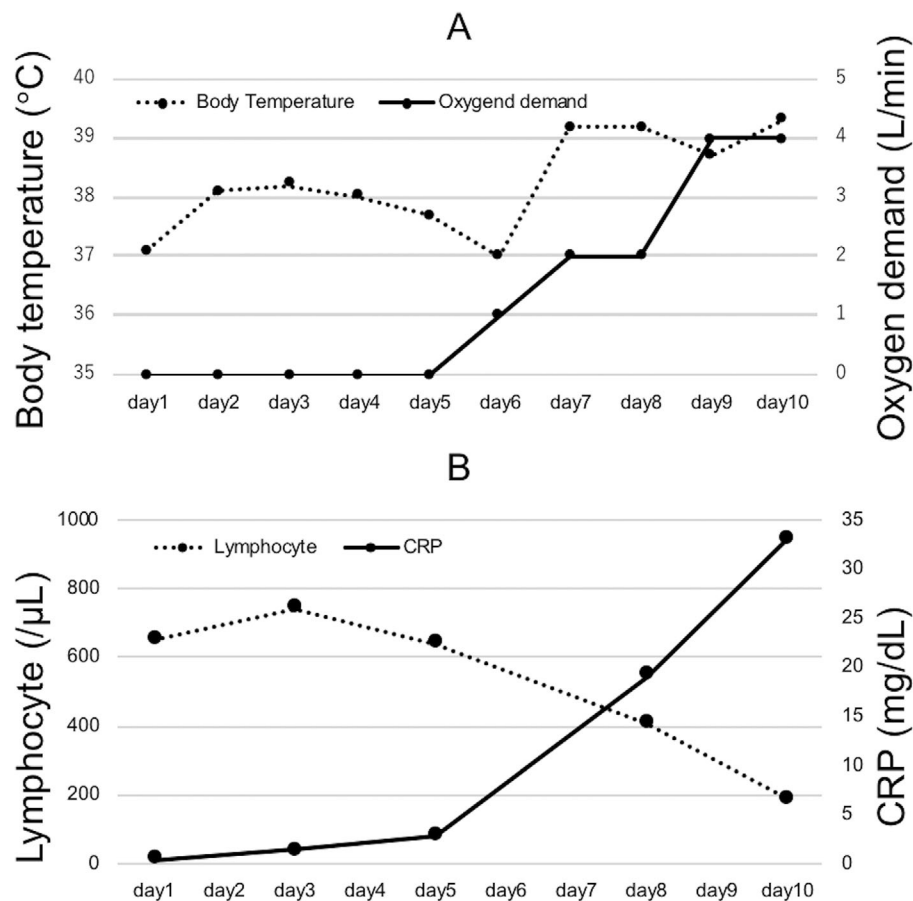
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**Figure 1.** Chest X-ray on hospitalization day 1 (A) and 8 (B). Slight, bilateral ground-glass opacity was evident on hospitalization day 1 and showed exacerbation on day 8.

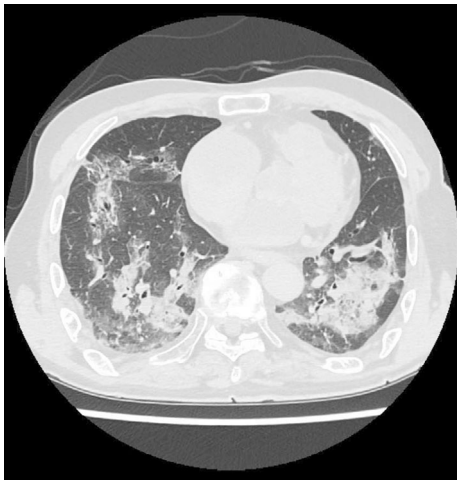


**Figure 2.** Lymphocytes decreased inversely to the exacerbation of the body temperature, hypoxemia, and CRP. The course of the body temperature and oxygen demand (A). The course of lymphocyte count and CRP levels (B).

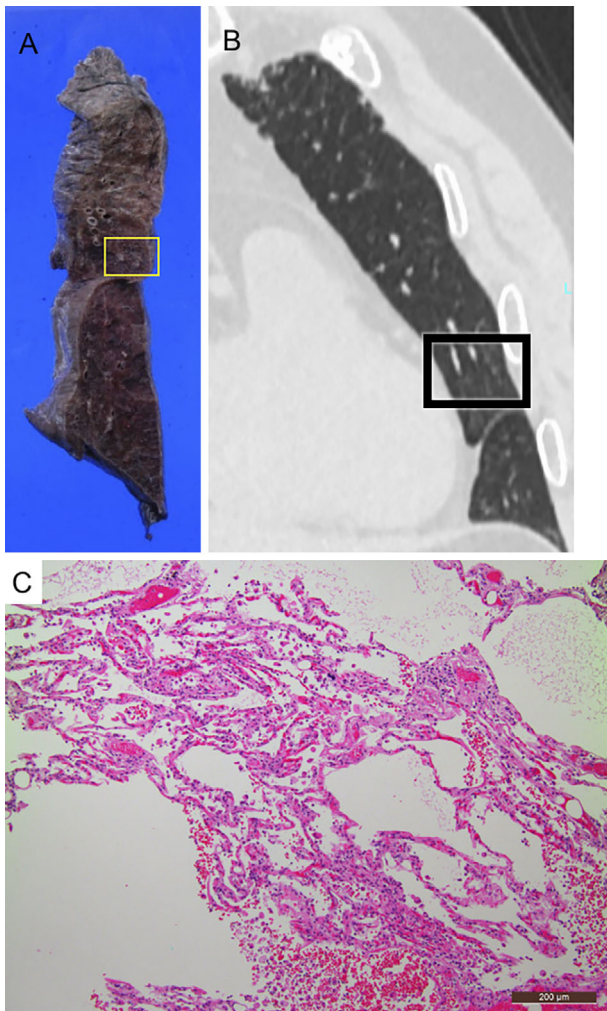
of 30 to 40 breaths per minute appeared. At 7:40, his pulse suddenly dropped from 105 beats per minute to cardiac arrest. Only four days had lapsed from the commencement of oxygen administration to death (Fig. 2A). Post-mortem computed tomography (CT) performed four hours after death showed extensive ground-glass opacities in the bilateral

lungs that were more severe in the lower dorsal lobes than in the upper lobes (Fig. 3), as the upper lobes of the lungs appeared relatively normal.

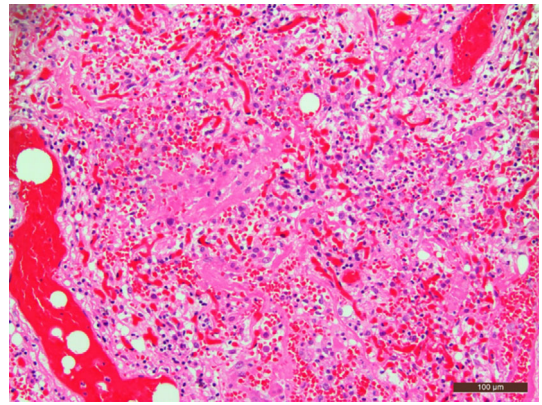
Because the patient's death had followed a relatively rapid clinical course, a pathological autopsy was performed to inquire into the direct cause of death. The autopsy of the body



**Figure 3.** Postmortem CT findings showing extensive bilateral ground-glass opacities in the lower dorsal region.



**Figure 5.** Lymphocytic infiltration confirmed in areas without abnormal findings on CT. Macroscopic findings (A), CT autopsy findings (B), and microscopic findings (C). The area shown in (C) corresponds to the area in the yellow (A) and black (B) boxes.



**Figure 4.** Pathological findings of the dorsal side of the lower lung lobe showing significant alveolar hemorrhaging.

(except for the brain) was conducted 22 hours after death. No pulmonary embolism or myocardial infarction was observed on a macroscopic examination. Congestion was systemically prominent. Petechiae were scattered on the lung surface. Microscopically, findings corresponding to the exudative phase of diffuse alveolar damage (DAD), including hyaline membrane formation, interstitial edema, lymphocyte infiltration, and type I alveolar epithelial cell shedding, were diffusely observed in the lungs. Significant alveolar hemorrhaging had occurred at the dorsal side of the lower lobe of the bilateral lungs (Fig. 4). Lymphocyte infiltration in the periphery of the blood vessels and edematous thickening of the blood vessel wall were observed. Lymphocytic infiltration into the interstitial tissue was also confirmed in areas where there were no abnormal findings on CT (Fig. 5). Based on the absence of pulmonary embolism and myocardial infarction, COVID-19 pneumonia was anatomically diagnosed as the cause of death.

## Discussion

Anti-inflammatory therapies are considered effective for the treatment of COVID-19 pneumonia. There are several clinical trials in which glucocorticoid therapy has shown good efficacy. In one randomized, open-label, clinical trial in Brazil enrolling 299 patients with confirmed or suspected COVID-19 who were receiving mechanical ventilation, intravenous dexamethasone plus standard care resulted in a statistically significant increase in the number of ventilator-free days compared with standard care alone (3). In another randomized open-label trial in the United Kingdom, oral or intravenous dexamethasone reduced the 28-day mortality among hospitalized patients to a greater extent than usual care alone (4). In that trial, which included hospitalized patients with COVID-19, 2,104 and 4,321 patients were randomized to a dexamethasone treatment group and a usual care group, respectively. As a result, 482 patients (22.9%) in the dexamethasone group and 1,110 patients (25.7%) in the usual care group died within 28 days after randomization (age-adjusted rate ratio, 0.83; 95% confidence interval [CI],



0.75 to 0.93;  $P < 0.001$ ) (4). In contrast, in a randomized, double-blind, sequential trial conducted in France that included 149 critically ill patients with biologically confirmed or suspected COVID-19 (based on chest CT findings in the absence of any other cause of pneumonia), low-dose hydrocortisone did not significantly reduce treatment failure (defined as death or persistent respiratory support) at day 21 compared with a placebo (5). Furthermore, the incidence of adverse events can increase even with short-term glucocorticoid administration (6). However, in a meta-analysis of 7 trials enrolling 1,703 critically ill patients with COVID-19, glucocorticoid therapy reduced mortality to a greater extent than standard care or a placebo (1). Based on these somewhat contradictory findings, it has been suggested that oxygen demand, severity of lung imaging findings, and risk factors, such as age and complications, should be considered when weighing the merits of using glucocorticoid therapy.

Lymphopenia is another risk factor of COVID-19 pneumonia aggravation (7). One hypothesis of the pathophysiology underlying COVID-19-associated lymphopenia is excessive recruitment and sequestration of lymphocytes in the lung (8). Our patient had lymphocytic infiltration in the interstitial tissue of the lungs, and his lymphopenia progressed as the disease progressed (Fig. 2B), supporting this hypothesis. Furthermore, lymphocytic infiltration was confirmed at sites that appeared normal on CT. Given the discrepancy between the imaging and histopathological findings, lymphocytic infiltration may have existed in a wider area than was indicated by radiographic findings.

Generally, glucocorticoids induce apoptosis of lymphoid cells (9); lymphocytic infiltration in the interstitial tissue in COVID-19 is thus the target of glucocorticoid therapy (10). In the present case, the DAD findings were the main histopathological change, and alveolar hemorrhaging was also observed. Therefore, it might have been difficult to save this patient even if glucocorticoid therapy had been started soon after the development of adult respiratory distress syndrome (ARDS). However, since the lymphopenia in this case preceded the rapid progression of the pulmonary lesions, early glucocorticoid administration might have conversely led to a better clinical course. Thus, in patients with a risk of severe exacerbation, lymphopenia can be a factor justifying early glucocorticoid administration, even if imaging studies reveal

only mild abnormalities.

**The authors state that they have no Conflict of Interest (COI).**

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**Patient consent:** Consent to publish the details of the case was obtained from the family of the patient.

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