An ongoing outbreak of a coronavirus disease 2019 (COVID-19) hit major cities of China, Wuhan, late December 2019 and subsequently spread to other areas of China and countries (1). As of February 16, 2020, there were 68,594 confirmed cases in mainland China. Among the confirmed cases, approximately 14.0% were severe (2). The mortality rate of severe patients was 6% (3). Herein, we described the clinical characteristics and chest CT follow-up of a severe COVID-19 patient, whose CT appearance was only moderate, and thus represents a case of mismatch between clinical severity and chest imaging severity.

A 39-year-old man presented to the hospital on Jan 24, 2020 with a week of fever, nonproductive cough and fatigue. He lived in Wuhan before onset, with no underlying diseases. On admission, his body temperature elevated to 38.6 °C (101 °F) with rough breathing sounds. Real-time fluorescence polymerase chain reaction (RT-PCR) of the patient’s throat swab was positive for 2019-nCoV nucleic acid. Laboratory findings indicated elevated C-reactive protein of 24.86 mg/L (normal rage, 0–6 mg/L), lactic dehydrogenase of 339 U/L (normal range, 120–250 U/L), and urine protein of 1+ g/L. White blood cell count (3.66×10⁹/L; normal range, 4.0–10.0×10⁹/L) and lymphocyte count (1.31×10⁹/L; normal range, 1.1–3.2×10⁹/L) were normal. Liver function, myocardial enzymes and coagulation function were normal. Electrocardiogram showed sinus tachycardia.

The initial unenhanced chest CT of Jan 24, 2020 showed bilateral group-glass opacities (GGO) located in subpleural regions (4,5) (Figure 1A). The chest CT was classified at early stage. However, the patient was diagnosed as severe according to clinical criteria. The patient was treated with antiviral drugs (oseltamivir and arbidol), anti-inflammatory drugs (e.g., moxifloxacin) and mask oxygen inhalation (flow rate, 2–4 L/min). Follow-up chest CT of Jan 28, 2020 showed GGO with consolidation and air bronchogram (progressive stage) (Figure 1B). Follow-up chest CT of Jan 30, 2020 showed reduced GGO with pulmonary fibrosis (absorption stage) (Figure 1C). The patient was then at clinically severe stage for the partial arterial oxygen pressure (PaO₂) and oxygen saturation (SO₂) was 65 mmHg, 92%, respectively. On Feb 01, the patient’s PaO₂ and SO₂ further decreased to 51 mmHg and 86%. The patient had no fever and cough. The repeat RT-PCR of Feb 18 was negative. The chest CT of Feb 19, 2020 showed obvious absorption of lesions (Figure 1D). The RT-PCR tests were repeated on Feb 24 and 25 and both were negative. On Feb 22, the patient’s SO₂ elevated to a normal value of 98%. All laboratory findings were normal. Therefore, the patient met the current criteria for hospital discharge (6) and left hospital in the followed week.

This case may be an example of a group of patients who are clinically diagnosed severe but at early-stage on chest CT at presentation. Their lung diseases do not rapidly progress more than 50% within 24–48 hours. The clinical outcomes of these patients are usually favorable. Hence,
Figure 1 A 39-year-old man living in Wuhan presenting with fever and cough. (A) Axial thin-section (1 mm) non-contrast chest CT image from Jan/24/2020 shows bilateral group-glass opacities (GGO) with a peripheral distribution; (B) follow-up chest CT image on Jan/28/2020 shows GGO with consolidation; (C) follow-up chest CT image on Jan/30/2020 shows GGO and consolidation absorption; (D) follow-up chest CT image on Feb/19/2020 shows further resolution.

the stage of these patients perhaps can be down-graded, especially for patients without underlying diseases, such as chronic obstructive pulmonary disease, hypertension, and diabetes. By contrast, a recent study showed symptoms relief but progression on chest CT, indicating that clinical symptoms and imaging findings are unmatched in early stage of COVID-19 (7). Therefore, the staging of COVID-19 should consider the severity on chest imaging, which will benefit for treatments and prognosis assessment.

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Footnote

Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at http://dx.doi.org/10.21037/qims.2020.03.12). The authors have no conflicts of interest to declare.

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