Serum amyloid A is a predictor for prognosis of COVID-19

To the Editors:

The coronavirus disease 2019 (COVID-19) outbreak originated in Wuhan, China, in December 2019. It has now affected 197 countries. By 27 April 2020, it was reported that 2,878,196 individuals were infected, with 6.1% of patients diagnosed with severe illness, and more than 198,000 deaths worldwide. Early identification and prediction of acute exacerbation is of great importance for the management of COVID-19. However, a serum biomarker for prognosis of COVID-19 is currently lacking.

Serum amyloid A (SAA) is commonly elevated in the acute phase of inflammatory diseases. We describe levels of serum SAA, C-reactive protein (CRP) and procalcitonin (PCT) in patients with COVID-19 during their hospital admission.

We recruited patients with COVID-19 from 20 January to 10 March 2020, and detected serum markers (SAA, CRP and PCT) within the first day of hospitalization. This study was approved by the Hospital Ethics Review Committee (Ethics No 20201134) and the patients’ informed consent was exempted. The outcomes of these patients were recorded as either improved and discharged or acutely exacerbated, according oxygenation status (oxygen saturation < 93% and arterial partial pressure of oxygen/oxygen concentration ≤ 300 mm Hg) and chest radiograph (>50% lesions progression within 24–48 h in pulmonary imaging). Logistic regression model and receiver operating characteristic (ROC) curve were analysed to investigate the possible roles of SAA, CRP and PCT in prognosis prediction of COVID-19.

The current observational study enrolled 118 patients with COVID-19 (64 males) whose average age was (mean ± SD) 49.55 ± 15.95 years and the time of illness onset ranged from 1 to 14 days. On admission, 16 cases were diagnosed as severe COVID-19 and the remaining 102 patients were identified as ordinary cases, who received further medical surveillance to investigate the prognosis. The levels of SAA, CRP and PCT were markedly elevated in patients with severe illness compared with ordinary cases (SAA (mean ± SD): 40.42 ± 52.62 vs 198.32 ± 55.12 mg/L, P < 0.001; CRP (mean ± SD): 16.55 ± 10.99 vs 46.52 ± 35.21 mg/L, P < 0.001; PCT (mean ± SD): 0.051 ± 0.041 vs 0.125 ± 0.148 ng/mL, P < 0.001) (Fig. 1). Furthermore, all 102 ordinary patients received antiviral therapy (Arbidol, Suzhou, China), of which 71 patients recovered and were discharged, but 31 cases underwent acute exacerbation. Logistic regression showed that SAA, but not CRP or PCT, could serve as an independent predictive factor of severe COVID-19, with an accuracy of 89.1% in predicting acute exacerbation (cut-off value: 122.9). Further evidence needs to be collected to confirm the possible correlation between SAA and the severity of outcome in COVID-19.

Figure 1 The level of inflammatory indicators in patients with COVID-19 on admission (●, will worsen; □, has worsened; ◆, remain mild). COVID-19, coronavirus disease 2019; CRP, C-reactive protein; PCT, procalcitonin; SAA, serum amyloid A.
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Key words: coronavirus, COVID-19, pneumonia, prognosis, serum amyloid A.

Author contributions: Conceptualization: X.-N.M., D.-F.C., L.S., H.-K.W., S.-Y.L. Data curation: D.-F.C., H.P., R.-C.C., L.S., H.-K.W. Formal analysis: X.-N.M., R.-C.C., L.S., H.-K.W. Investigation: H. P., R.-C.C., L.S., H.-K.W. Methodology: X.-N.M., D.-F.C., R.-C.C., L. S. Project administration: X.-N.M., C.-L.L. Resources: X.-N.M., C.- L.L., H.P., S.-Y.L. Supervision: C.-L.L., S.-Y.L. Validation: C.-L.L., S.-Y.L. Writing—original draft: X.-N.M., C.-L.L., D.-F.C. Writing—review and editing: X.-N.M., C.-L.L., D.-F.C., S.-Y.L.

Abbreviations: AUC, area under the curve; COVID-19, coronavirus disease 2019; CRP, C-reactive protein; OR, odds ratio; PCT, procalcitonin; ROC, receiver operating characteristic; SAA, serum amyloid A.

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