Multisystem Inflammatory Syndrome Related to COVID-19 in Children

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Pediatric inflammatory multisystem syndrome temporally associated with severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (PIMS-TS) is a rare but serious complication of coronavirus disease 2019 (COVID-19). Reports of PIMS-TS started appearing around the world in the months after the pandemic began [1].

Two recent studies totaling 75 pediatric patients published in JAMA showed that PIMS-TS is novel and different from Kawasaki disease (KD) and toxic shock syndrome (TSS) [2, 3].


In the study by Whittaker et al [2] patients’ clinical features were compared with those of children diagnosed as having KD or KD shock syndrome [4]. Forty-five of 58 patients (78%) had evidence of current or previous infection with the SARS-CoV-2 that causes COVID-19. Thirty-three (57%) were females, 12 were white, and all were previously healthy, with mild asthma in three, and their median hospital stay was 7.1 days.

All 58 patients had fever, 31 (53%) had abdominal pain, 30 (52%) had a rash, 26 (45%) had conjunctivitis, 29 (50%) developed shock and needed supportive drugs or fluids, 17 (29%) had mucus membrane changes and cracked lips, 15 (26%) had headache, nine (16%) had enlarged lymph nodes, 13 (22%) had kidney damage, nine (16%) had swollen hands and feet, eight (14%) had dilated coronary arteries or aneurysms, and six (10%) had a sore throat. Four patients (7%) developed abnormal heart rhythms. Twenty-five (43%) required mechanical ventilation, and two (3%) needed extracorporeal membrane oxygenation for severe heart dysfunction.

Thirteen individuals met the American Heart Association criteria for KD [5], and 23 had fever and inflammation without characteristics of shock or KD. When compared with children with KD and TSS, PIMS-TS patients were older (median age: 9 vs. 2.7 years in KD and 3.8 years in TSS), and had elevated levels of C-reactive protein.

Whittaker et al [2] identified three unique features for presentations of PIMS-TS: 1) Persistent fever and high levels of inflammatory markers without features of KD, shock, or organ failure; 2) Characteristic signs and symptoms of KD; 3) Shock and clinical, echocardiographic, and lab evidence of heart damage.

They identified a pattern of cytokine expression that suggested an interferon signaling component, along with interleukin (IL)-6 and IL-10 production, similar to the one observed in KD and acute pulmonary COVID-19 infection. They concluded that the absence of elevated tumor necrosis factor (TNF)-α or IL-13 levels may differ from acute pulmonary COVID-19 infections [6].

All 17 patients in the study by Cheung et al [3] had a fever (median of 5 days), 14 had gastrointestinal symptoms, 12 had a rash, 11 had conjunctivitis, and nine had red, swollen lips. Three had low oxygen levels, and 13 went into shock. Fourteen had abnormal chest radiograph findings, and eight met the criteria for KD and five for incomplete KD [5].

All children had elevated levels of inflammatory markers, 16 had high serum IL-6 levels, 15 had high levels of N-terminal-pro-B-type natriuretic peptide (NT-proBNP), 14 had high levels of troponin T, 12 had reduced levels of lymphocyte white blood cells, and 11 had high levels of the immature white band cells.

Fifteen children required intensive care, and 10 needed support for low blood pressure. Nine patients had low oxygen levels, but none required mechanical ventilation. Three patients had abnormal heart rhythms, and one developed a medium-sized aneurysm.

Median patient age was 8 years (range: 1 - 16 years), nine were females, 12 were white, and all were previously healthy, with mild asthma in three, and their median hospital stay was 7.1 days.

The findings of these two studies can help clinicians characterize the clinical features of hospitalized, seriously ill children with PIMS-TS and provide insights into this apparently novel syndrome. The occurrence of abnormal cardiac findings suggests the need for long-term surveillance. Early recognition of PIMS-TS would enable pediatrician provide close monitoring and the optimal treatment.
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Financial Disclosure

None to declare.

Conflict of Interest

None to declare.

Data Availability

The author declares that data supporting the findings of this study are available within the article.

References