SARS-CoV-2-related Multisystem Inflammatory Syndrome in Adult complicated by myocarditis and cardiogenic shock

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Abstract

Multisystem Inflammatory Syndrome in Adult (MIS-A) is a rare COVID-19 complication, presenting as fever with laboratory evidence of inflammation, severe illness requiring hospitalization and multisystem organ involvement. We report on a 25-year-old man presenting with fever, rash, abdominal pain, diarrhoea and vomiting following prior asymptomatic COVID-19 infection. He developed refractory shock and type 1 respiratory insufficiency requiring mechanical ventilation. Diagnostic testing revealed significant inflammation, anemia, thrombocytopenia, acute kidney injury, hepatosplenomegaly, colitis, lymphadenopathy and myocarditis necessitating inotropy. Ventilatory, vasopressor and inotropic support was weaned following pulse corticosteroids and intravenous immunoglobulins. Heart failure therapy was started. Short-term follow-up shows resolution of inflammation and cardiac dysfunction.

Keywords
Acute heart failure; Myocarditis; Covid-19; echocardiography; cardiac MRI; intensive care medicine

Introduction

Severe Acute Respiratory Syndrome CoronaVirus 2 (SARS-CoV-2) is the causative agent of the current CORonaVIrus Disease-2019 (COVID-19) pandemic. About 40% of adult COVID-19 infections are asymptomatic. Clinical presentation of COVID-19 is heterogeneous ranging from a mild to severe respiratory phenotype. Gastro-intestinal and dermatological symptoms are also reported, suggesting multisystem involvement. Some patients present with critical respiratory disease, often complicated by acute respiratory distress syndrome, acute kidney injury, thromboembolism and secondary infections with septic shock, associated with high mortality. In children, clinical presentation is generally mild and rarely critical. An atypical Kawasaki disease or toxic shock-like syndrome was reported, occurring four weeks after an asymptomatic or mild COVID-19 infection in children and adolescents up to 21 years old, termed Multisystem Inflammatory Syndrome (MIS-C). Rare reports of this syndrome were described in adults (MIS-A).

Case Report

A 25-year-old man, with a recent (four weeks earlier) asymptomatic COVID-19 infection, was referred to our Intensive Care Unit (ICU) for evaluation and treatment of an unexplained shock syndrome. There was no history of smoking, alcohol or drug use. There was no exposure to other infectious or toxic agents and he exercised recreationally. The patient was double BNT162b2 vaccinated with the last dose nine months before admission. He presented to his general practitioner with fever (>39°C), abdominal pain, vomiting
and diarrhoea for four days. Ambulatory biochemical evaluation revealed elevated Erythrocyte Sedimentation Rate and C-Reactive Protein (CRP) with normal white blood cell count (Supplementary Table S1). Chest radiography was normal (Figure 1A). Amoxicillin/clavulanic acid was prescribed which was switched to cefuroxime following the occurrence of a diffuse erythematous and maculopapular rash.

Worsening clinical condition caused the patient to present to the emergency department. Clinical examination showed an ill patient with dyspnoea and diffuse abdominal pain. Heart-lung auscultation was normal. Rash persisted with associated bilateral conjunctivitis without other mucosal lesions. Blood pressure was 141/80 mmHg, pulse 110 bpm, temperature 37.8°C and oxygen saturation 98% without supplemental oxygen. ECG showed sinus tachycardia with new, diffuse PR-depression (Figure 2). Blood analysis demonstrated neutrophilia, thrombocytopenia, lymphopenia, high ferritin and D-dimer test, CRP doubling since visiting his GP and increased serum creatinine, gamma-glutamyl transferase, alkaline phosphatase and total bilirubin (Supplementary Table S1). Thoraco-abdominal Computed Tomography (CT) was performed because of suspected intra-abdominal sepsis and revealed uncomplicated ileocaecal inflammation, mesenteric lymphadenopathy, splenomegaly, normal lung parenchyma and discrete pericardial effusion. Transthoracic echocardiography (TTE) showed normal cardiac morphology and function without significant pericardial effusion. Blood, sputum, stool and urine cultures were incubated and the patient was switched to intravenous ciprofloxacin and metronidazole because of suspected intra-abdominal sepsis. Over the next two days, the patient developed hypotension and high anion gap metabolic acidosis due to lactic acid accumulation, necessitating fluid resuscitation and norepinephrine. Concurrently, he developed type 1 respiratory insufficiency treated with High Flow Nasal Oxygen and, finally, sedation, intubation and mechanical ventilation.

Because of further clinical deterioration, thoraco-abdominal CT was repeated upon arrival in our hospital and showed terminal ileitis and diffuse colitis with increased lymphadenopathy, hepatosplenomegaly, new ascites and bilateral pleural effusions (Figure 3). Failure of shock resolution and persistent inflammation prompted antimicrobial escalation to meropenem and vancomycin. With persistent suspicion for intra-abdominal septic shock and no etiological explanation on repeat imaging, an urgent exploratory laparoscopy was performed and showed moderate clear ascites without any signs of infection. Ascites was recovered for culture. Because of unexplained shock with increasing vasopressor doses, worsening peripheral perfusion and increasing lactate, a repeat TTE was performed. It showed a mildly dilated left ventricle with moderately decreased ejection fraction due to anteroseptal akinesia and hypokinesia of the other segments, as well as a 6 mm circumferential pericardial effusion without tamponade physiology (Supplementary Video S1 and Supplementary Table S2). New chest radiograph showed cardiomegaly, bilateral pleural effusions and pulmonary edema (Figure 1B). High sensitive troponin T was dynamically elevated up to 288 ng/L and NT-pro-BNP was 28,770 pg/mL (Supplementary Table S1). Because of suspected myocarditis and cardiogenic shock, dobutamine was associated, as well as levosimendan. Cardiac MRI (CMR) was performed four days after hospital admission and showed a moderately dilated left ventricle with moderately reduced systolic function, evidence for non-ischemic myocardial injury (increased myocardial extracellular volume by T1-mapping) and focal myocardial edema in the anterior and apicolateral segments on T2-weighted images, but no focal myocardial late gadolinium enhancement, fulfilling the 2018 Lake Louise criteria for myocarditis (Figure 4).16 Pericardial inflammation was
apparent by the presence of pericardial effusion, thickening and gadolinium enhancement on CMR (Figure 4). Retrospective CT evaluation confirmed normal coronary arteries. Rhythm monitoring revealed no arrhythmia. Cultures, auto-immune and viral serology remained negative. PCR revealed persistently high SARS-CoV-2 viral load, more than four weeks after the initial positive test, in the presence of anti-N IgG.

MIS-A was diagnosed because of the suggestive rash, conjunctivitis, gastro-intestinal symptoms, thrombocytopenia, fever, major inflammation, shock and severe myocarditis with cardiac failure and lack of any other explanation and underlying infection. Intravenous immunoglobulins (IVIGs, 2 g/kg for 1 day, maximum 100 g) and pulse corticosteroids (10 mg/kg/d for 3 days, maximum 1 g/d) were started. The patient improved and was weaned off ventilatory,

![Figure 2](image1.png)

**Figure 2** 12-lead ECG showing sinus tachycardia and diffuse PR-depression. ECG recorded at 25 mm/s and 10 mm/mV.

![Figure 3](image2.png)

**Figure 3** Computed tomography following transfer to the academic ICU department showing bilateral pleural effusion and mild pericardial effusion (white arrowheads) (panel A), hepatosplenomegaly (panel B) and colitis with adjacent ascites (white arrowheads) in the right lower quadrant (panel C).
inotropic and vasopressor support within three days. Antibiotics were discontinued two days later. ECG showed regression of PR depression. TTE showed improved anteroseptal contractility, increased LVEF and regressed pericardial effusion (Supplementary Table S2). Corticosteroids were continued orally (2 mg/kg/d) and slowly tapered over six weeks. Proton pump inhibitor and calcium/vitamin D supplements were associated. Heart failure therapy (bisoprolol 2.5 mg and lisinopril 10 mg daily) was titrated. Because left ventricular function improved rapidly following start of pulse corticosteroids and IVIGs, lisinopril was not escalated to valsartan-sacubitril and no mineralocorticoid receptor antagonist was associated. Low-dose aspirin was associated to prevent vascular events. At ambulatory short-term follow-up following hospital discharge the patient was asymptomatic and without inflammation. Resting ECG was normal and TTE showed lower septum and posterior wall thicknesses, as well as normalisation of left ventricular diameters and systolic function (Supplementary Table S2 and Supplementary Video S2). Heart failure therapy and low dose aspirin will be discontinued in the future.

**Discussion**

COVID-19-associated MIS was first reported in children in April 2020 as a Kawasaki disease- and toxic shock-like hyperinflammatory syndrome. The Center for Disease

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**Figure 4** Cardiac MRI scan showing dilated left ventricle and mild pericardial effusion (white arrowheads) on cine images (panels A). Myocardial edema in the anterior and lateral apical segments (white arrowheads) on STIR-T2 weighted images (panels B). Pericardial enhancement, and no myocardial late gadolinium enhancement (panels C). Pre- and postcontrast T1 maps from which a myocardial extracellular volume of 31% (normally 25.3 ± 3.5%) was calculated (panels D). Indexed myocardial mass was augmented (193 g/m², normally <107 g/m²).

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Control (CDC) released a MIS-C definition (Box 1) in May 2020. Since June 2020, a similar syndrome was described in adult patients. MIS is a rare COVID-19 complication: 316 per 100,000 SARS-CoV-2 infections in children. Currently, more than 221 MIS-A cases have been reported and analysed in two systematic reviews. Patients generally develop MIS-A four weeks after COVID-19 infection. Most are male (70%) and 21–34 years old, but the oldest is 84 years. Currently, COVID-19 vaccination is not associated with MIS.

MIS-A is diagnosed by exclusion. The CDC defines MIS-A in patients aged ≥ 21 years old hospitalized for ≥ 24 hours, or with an illness resulting in death, who meet the clinical criteria and laboratory evidence summarized in Box 1. The criteria must be met by the end of the third hospital day. In this case, intra-abdominal sepsis was initially suspected and repeated cultures, assessments by computed tomography and a laparoscopy were performed to exclude sepsis before initiating immunomodulatory treatment for MIS-A. In this 25-year old man, MIS-A was diagnosed at the start of the third hospital day based on the presence of two primary (severe cardiac illness in combination with rash and non-purulent conjunctivitis) and three secondary clinical criteria (shock, abdominal pain and thrombocytopenia) in the presence of elevated CRP and ferritin as well as recent positive SARS-CoV-2 PCR.

Although not mandatory for the diagnosis of MIS-A, endomyocardial biopsy (EMB) may be performed in suspected myocarditis and may provide differentiation between inflammatory, infectious and infiltrative causes. In MIS-A, EMB histopathology has shown an inflammatory infiltrate of macrophages, T lymphocytes and eosinophils. It may be particularly useful if histological evaluation is expected to impact therapy. In this case, we estimated EMB would not add diagnostic clues given the typical clinical course of MIS-A in the absence of a more likely alternative diagnosis, nor would it alter our therapeutic management. And indeed, the patient improved rapidly upon treatment with pulse corticosteroids and IVIGs.

The organ systems most affected in MIS-A are hematological (41.8–92%), cardiovascular (81–87%), gastro-intestinal (73.4–83%) and respiratory (29.1–74%). Musculoskeletal (46–52.1%), neurological (16.4–47%) and renal (43%) manifestations are...
less frequent.\textsuperscript{22,23} Organ system involvement is comparable between adults and children.\textsuperscript{31} Cardiovascular findings in MIS-C/A include perimyocarditis (11–33\%), hypotension and shock (50–60\%), coronary artery dilatation or aneurysm (4–8\%), mitral regurgitation (14\%) and arrhythmia (12–18\%).\textsuperscript{22,23,31–33} Left ventricular dysfunction is present in 55\% of adults with mean left ventricular ejection fraction (LVEF) 39\%.\textsuperscript{22,23} Left ventricular dysfunction with ejection fraction <55\% is present in 38\% of MIS-C patients.\textsuperscript{31–33} Additionally, 28.6\% of MIS-A patients and 3\% of MIS-C patients show severe LV dysfunction with LVEF <30\%.\textsuperscript{22,23,31–33} Patients with LVEF <45\% are more likely to present in shock (60\% vs 9\%) and be admitted to ICU (100\% vs 55\%) as compared to patients with LVEF ≥45\%.\textsuperscript{34} Patients with lower LVEF also resided longer on ICU with more need for inotropy, but showed no significant differences in total hospital stay, ventilatory support or mortality. Treatment generally causes normalisation of LVEF after 1 week.\textsuperscript{22,23,31–33} Mortality is 1.9\% in MIS-C and 7\% in MIS-A and has been reported in patients that require mechanical circulatory support and do not show improvement in LVEF.\textsuperscript{22,35}

Early diagnosis is crucial since MIS-C/A is severe, but treatable. Cardiovascular complications triggered recommendations for immunomodulation and intensive cardiac observation.\textsuperscript{34,36} Cardiac complications generally regress with these interventions.\textsuperscript{22,23,31–33} Cardiac involvement may be diagnosed by a combination of ECG, cardiac biomarkers, echocardiography and radiography. Very limited data (9 adults, 4 children) are available on CMR in MIS.\textsuperscript{37,38} This case report shows that the 2018 Lake Louise criteria may be performant for the diagnosis of MIS-associated myocardial inflammation. More research is needed to describe long term effects and identify risk factors or imaging criteria for MIS development and associated mortality. MIS pathophysiology is currently incompletely understood. An immunopathological cascade is suggested based on natural history, immunological phenotyping and histopathology.\textsuperscript{39–43} Failure of the innate and adaptive immune response to SARS-CoV-2 causes hyperinflammation. Abnormal IFNγ production activates macrophages, natural killer cells and T-cells, activating inflammatory cytokine cascades that damage multiple organ systems (Figure 5). In MIS-C, several studies have shown activation of oligoclonally expanded T cells, reminiscent of disease driven by superantigen exposure such as in toxic shock syndrome. These T cells harbor the \textit{TRBV11}–2 gene, encoding for the T cell receptor Vβ 21.3, which seems to be a highly sensitive and specific feature.\textsuperscript{39,43,44} The resemblance with Kawasaki disease and toxic shock syndrome spurred experts to adopt analogous treatments including IVIGs, aspirin, corticosteroids, and anti-IL-1, -IL-6, or -TNFα.\textsuperscript{37,45} Evidence-based data remain scarce to date.

Box 1. CDC MIS-C/A definition.

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<thead>
<tr>
<th>MIS-C</th>
<th>MIS-A</th>
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<tr>
<td>I. Clinical Criteria</td>
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<td>Subjective fever or documented fever (&gt;38.0 C) for ≥24 hours prior to hospitalization or within the first THREE days of hospitalization* and at least THREE of the following clinical criteria occurring prior to hospitalization or within the first THREE days of hospitalization*. At least ONE must be a primary clinical criterion.</td>
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<tr>
<td>Primary clinical criteria</td>
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<td>1 Severe cardiac illness includes myocarditis, pericarditis, coronary artery dilatation/aneurysm, or new-onset right or left ventricular dysfunction (LVEF&lt;50%), 2nd/3rd degree A-V block, or ventricular tachycardia. (Note: cardiac arrest alone does not meet this criterion)</td>
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<td>2 Rash AND non-purulent conjunctivitis</td>
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<td>Secondary clinical criteria</td>
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<tr>
<td>1 New-onset neurologic signs and symptoms includes encephalopathy in a patient without prior cognitive impairment,</td>
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(Continues)
MIS-C | MIS-A
---|---
seizures, meningeal signs, or peripheral neuropathy (including Guillain-Barré syndrome) | 
2 Shock or hypotension not attributable to medical therapy (e.g., sedation, renal replacement therapy) | 
3 Abdominal pain, vomiting, or diarrhea | 
4 Thrombocytopenia (platelet count <150,000/microliter) | 

II. Laboratory evidence The presence of laboratory evidence of inflammation AND SARS-CoV-2 infection. 
1 Elevated levels of at least TWO of the following: C-reactive protein, ferritin, IL-6, erythrocyte sedimentation rate, procalcitonin | 
2 A positive SARS-CoV-2 test for current or recent infection by RT-PCR, serology, or antigen detection | 

*Fever >38.0 °C for ≥24 hours, or report of subjective fever lasting ≥24 hours 
**Including, but not limited to, one or more of the following: an elevated C-reactive protein (CRP), erythrocyte sedimentation rate (ESR), fibrinogen, procalcitonin, d-dimer, ferritin, lactic acid dehydrogenase (LDH), or interleukin 6 (IL-6), elevated neutrophils, reduced lymphocytes and low albumin

*These criteria must be met by the end of hospital day 3, where the date of hospital admission is hospital day 0.

Acknowledgements

Figure 5 was constructed using open source clipart from https://smart.servier.com/.

Conflict of interest

The authors have no conflicts of interest to disclose.

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None with respect to the current manuscript.

Supporting information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Table S1. Results of laboratory testing.

Table S2. Longitudinal analysis of transthoracic echocardiography.
IVSd = end-diastolic Interventricular Septum thickness, LVDd = Left Ventricular end-diastolic Diameter, LVPWd = end-diastolic Left Ventricular Posterior Wall thickness, LVDs = Left Ventricular end-systolic Diameter, LVEF = Left Ventricular Ejection Fraction, RWMA = Regional Wall Motion Abnormalities, LVCO = Left Ventricular Cardiac output, LA = Left Atrium, RA = Right atrium, DT = Deceleration Time, RVEDD = Right Ventricular End-Diastolic Diameter, TAPSE = Tricuspid Annular Plane Systolic Excursion, sPAP = systolic Pulmonary Artery Pressure, CVP = Central Venous Pressure.

Video S1. Parasternal short axis view from transthoracic echocardiography following transfer to academic ICU department.
Video S2. Parasternal short axis view from transthoracic echocardiography at one week follow-up.
References


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