

A Rare Case of 10-year-old Child with Multisystem Inflammatory Syndrome (MIS-C)

Case Study

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Abstract

10-year-old male presented with 5 days of fever accompanied with cough, vomiting, muscle weakness, conjunctival injection, tachycardia and pharyngeal erythema alert but having toxic look. Detailed laboratory investigations revealed MIS-C with no covid contact history and RT PCR were negative. Dobutamine and methylprednisolone led to improved therapeutic outcome.

Keywords: MIS-C; Pediatric Patient; Post Covid Illness.

Introduction

The corona virus disease 2019 (Covid-19) pandemic has caused catastrophic disease worldwide. Children and infants have been relatively spared from severe COVID-19, with pediatric cases accounting for less than 2% of COVID hospitalizations. The U.S. Centre for Disease Control and Prevention (CDC) recently recognized a new syndrome in children and teens ages 2–15 that is associated with the coronavirus that causes COVID-19. While the syndrome is very rare, it can be dangerous. Multisystem inflammatory syndrome in children (MIS-C) was first identified in April 2020, by doctors at children's hospitals in the United States and the United Kingdom. The condition has also been called pediatric inflammatory multisystem syndrome (PIMS) [1, 2]. No such case has been reported from India till date. Here we report first case of MIS-C in paediatric patients from India.

Case Study

A 10-year-old previously healthy boy was presented with fever for 5 days with symptoms of cough, abdominal pain, vomiting, rash and conjunctivitis, fatigue and irritability. Vital parameters showed temperature 102.3°F, SpO₂-98%, Respirations-24/min and Heart rate-136/min. Physical examination demonstrated conjunctival redness, oedema, moist mucous membranes with

pharyngeal erythema without exudates, respiratory system clear to auscultation bilaterally, cardiovascular system: S1, S2 normal but tachycardia, abdomen was soft and non-tender (figure 1). Investigations revealed raised WBC count 9800 (Neutrophils-81, Lymphocytes-10, Eosinophils-4, Monocytes-4, Basophils-1) with haemoglobin 11.5g/dl and platelet count 1.66 lakhs. No covid positive contact history and X-ray chest was normal. A negative result was obtained for rapid antigen test for covid 19. Patient was admitted and every two hourly temperature was recorded. The patient continued to have spiking fever every 4-6 hours and tachycardia. Sleeping heart rate was 126 bpm without fever. The patient developed loose motion and abdominal pain. Preliminary diagnosis of patient lead to initial treatment of patient with therapy for mitigating viral infection using a moxycillin + clavulanic acid and Paracetamol accompanied with IV fluids (0.45% Dextrose with K+@1mEq/kg/day). Vitamin C, Zinc and Vitamin D Supplementation were also given to patient treat fatigue but unfortunately there was no significant improvement observed in the condition of patient.

Due to persistent high-grade fever (102.4°F) and tachycardia (146 bpm), the patient was shifted to paediatric ICU. Repeat investigations showed Hb 10.5gm%, Asignificant rise in TLC count (13000), neutrophils-80%, lymphocyte-11%, eosinophil-4%, monocyte-4%, basophils-1%, platelet 142000, CRP level (106.80

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Received: August 13, 2020

Accepted: October 07, 2020

Published: November 19, 2020

Citation: Ashish Goti, Ramesh Dihora, Zankhan Mirani, Manthan Mirani. A Rare Case of 10-year-old Child with Multisystem Inflammatory Syndrome (MIS-C). *Int J Pediatr Health Care Adv.* 2020;6(1):99-100. doi: <http://dx.doi.org/10.19070/2572-7354-2000028>

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Figure 1. Clinical presentation of the MIS -C.



mg/dl), D-dimer(1.2 mg/dl) and ferritin(284.82 ng/dl) was observed on 7th day of initial fever. Peripheral smear-negative for malarial parasite. The initial treatment was discontinued due to severe tachycardia (151bpm) and hypotension (76/50). 2D echo of the patient showed poor myocardial function(EF 30%). Patient was shifted to dobutamine therapy and suspected for MIS-C due to constant hypotension and decrease cardiac function. Hence immediately patient was tested for CovidIgG test which showed a positive result. The absence of preceding symptoms of COVID-19 indicated by negative polymerase chain reaction result with positive antibody test confirmed MIS-C. Immediately the patient was shifted on MIS-C therapy starting with IVIG(2g/kg), methylprednisolone(30mg/m²/day), aspirin(6-mg/kg/day), pantoprazole(40mg/day) and ceftriaxone (75 mg/kg IV)treatment.

Following the first dose of treatment congestion subsided along with improved heart function (EF 45%) with reduction in fever. On three days of continued treatment patient was shifted to paediatric ward prescribed with oral prednisolone therapy. The patient was discharged from the hospital on the 7th day of admission. The vital parameters of the patient were normal on discharge and the patient showed no symptoms on regular follow-up for two weeks.

Discussion

MIS-C or PIMS has features in common with toxic shock syndrome and an illness called Kawasaki disease, which cause inflammation of the blood vessels throughout the body. The association with this syndrome and COVID-19 may be due to the body's immune response to the presence of the SARS-CoV-2. Persistent symptoms of conjunctival redness, pharyngeal erythema with prolonged fever and chills, rash, myocarditis, elevated C-reactive protein level supported by IgG Antibody test revealed confirmation of MIS-C infection in patient with prior infection due to

COVID-19 virus which was in contrast to Kawasaki Disease [1]. The distinct symptoms demonstrating cardiac dysfunction or depression, coagulopathy, gastrointestinal symptoms accompanied by mild respiratory symptoms and occasional indications for supplemental oxygen was characteristic of MIS-C which was found to be a contrasting with most cases of acute Covid-19 among hospitalized children [2] and no complications related to circulatory failure in acute phase, artery aneurysm or resistance to intravenous immunoglobulin were observed as in Kawasaki Disease [3]. Diffuse myocardial edema and hyperaemia without evidences of focal myocardial necrosis which was in contrast to recent publications of myocarditis associated with COVID-19 [4] followed by treatment with IVIG, glucocorticoids, and vasopressors as immediate functional therapy was in consistent with other studies published [5]. Early diagnosis of cardiac and gastrointestinal symptoms followed by serial measurement of cardiac function with prompt treatment acts as a key to prevent multi-organ failure and long term organ damage due to MIS-C.

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