



COVID-19 CLINICAL ROUNDS: PEDIATRIC CRITICAL CARE

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Disclosure

- I have nothing to disclose.

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Pediatric Critical Care: COVID-19 and MIS-C

- ICU level cases of SARS-CoV-2 infection—May 2020
- Steady stream of admissions with peak PICU census of 7-8 patients/daily now declining
- Creation of a COVID-19 pod within our ICU
- 3-4 weeks later: Multi-System Inflammatory Syndrome in Children (MIS-C)
- Presentation is extremely variable
 - 2 cases today demonstrate the range of disease
- Children with both illnesses continue to trickle in

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Case #1: 5-year old male

- Previously completely healthy, except 34-week prematurity
- Presented with:
 - 5-day history of fever and fatigue
 - Diagnosed with strep pharyngitis by PCP-started on Amoxicillin
 - Mild abdominal pain and vomiting
 - Rash, eye redness, cracked lips
 - Chronically enlarged cervical nodes
 - No sick contacts, no h/o exposure to COVID
- Admitted to general ward service but transferred to PICU after 12 hours with shock

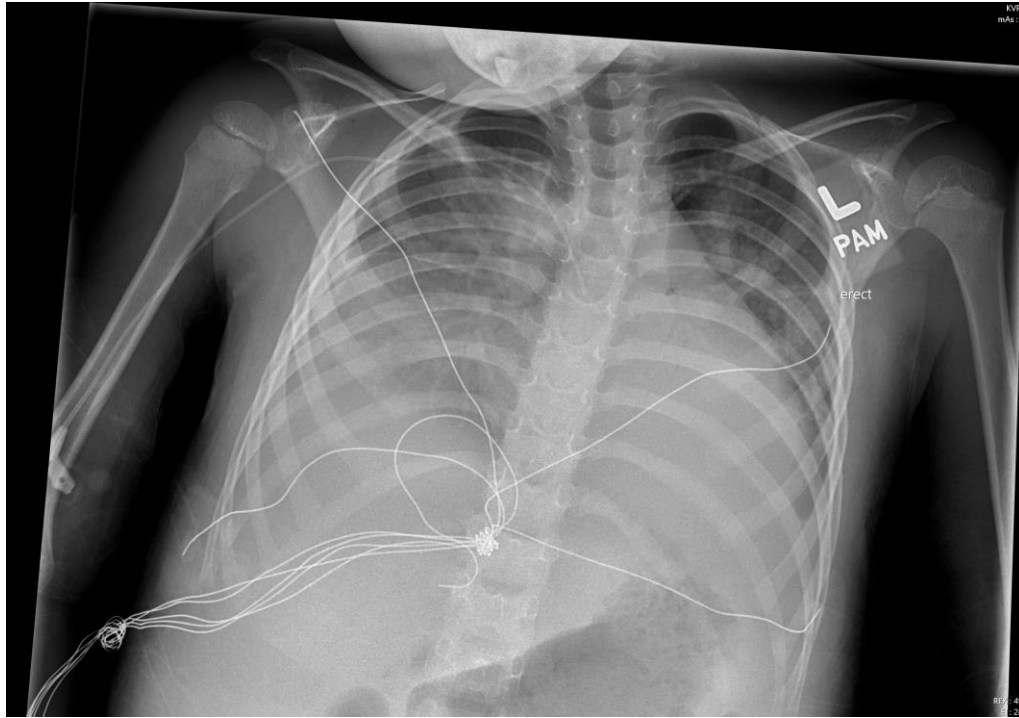
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Case, cont'd.

- Before arriving in the PICU, 100cc/kg of fluid resuscitation
- On epinephrine and norepinephrine infusions
 - ECHO without coronary dilation/aneurysm
- Acute kidney injury and oliguria
- Persistent RUQ pain—U/S shows hydrops of the gallbladder
- Labs suggest MIS-C: elevated BNP, elevated D-dimer and ferritin, high CRP and ESR, low platelets

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Acute respiratory failure on day 2-3 of hospital course



- Progressive respiratory insufficiency
- Managed with non-invasive positive pressure support
- Pleural effusion and inc. vascularity

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Case 1: diagnosis & treatment

- Was this Kawasaki or MIS-C??
 - COVID-19 antibody test (IgG) was positive
- Broad empiric antibiotic coverage
 - All cultures remained negative
- Treated with high dose steroids, IVIG
- Aggressive diuresis after shock reversed
- Enoxaparin for venous thrombosis prophylaxis—
aspirin at discharge
- Follow up with cardiology, rheumatology,
hematology

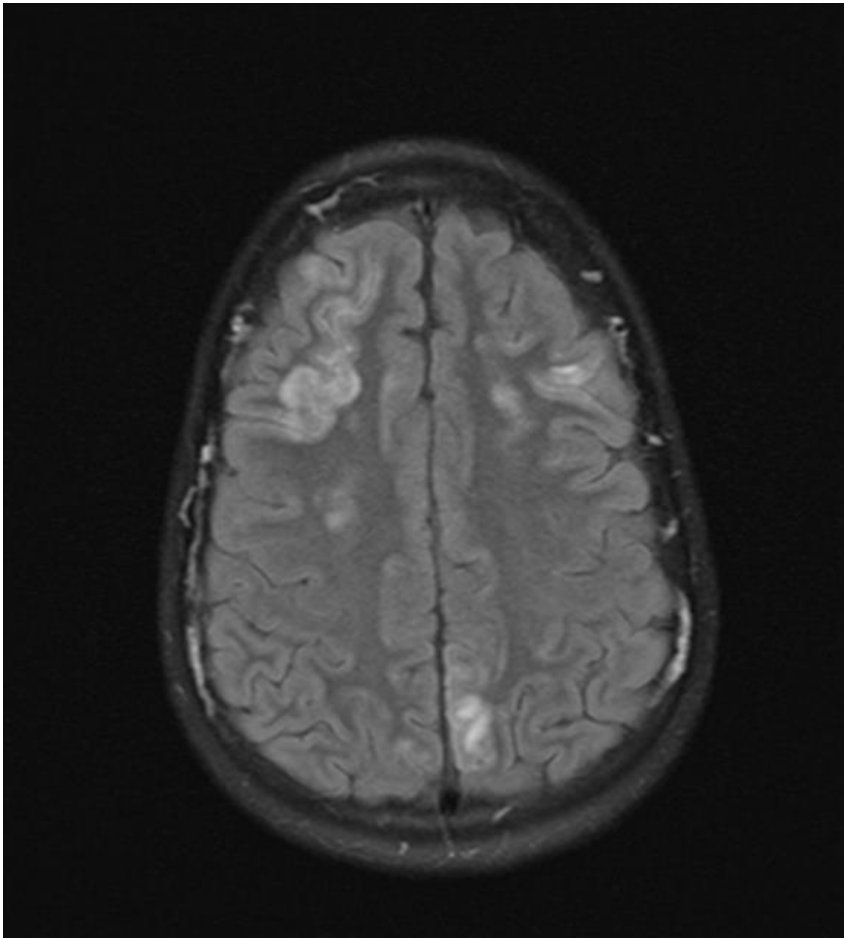


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Case 2: 11-year old male

- 11yo previously healthy male admitted 1 week after uncomplicated SARS-CoV-2 infection
- High fever → 105
- Cardiogenic shock-ECHO with depressed function and coronary artery dilation
- Acute respiratory failure
- 11-day hospital course with clear signs of MIS-C
 - Elevated ESR, CRP, D-dimer + all organ injury
- Treated with steroids and IVIG and enoxaparin
- Returns 2 weeks later with abdominal pain, vomiting, r/o appendicitis
- Neurologic decompensation night 1—onset of seizures
 - Emergent intubation, brief CPR, transfer to ICU

MRI abnormalities



- Scattered foci of cortical edema in bilateral frontal, right temporal, left parietal lobes-PRES?
- But only mild hypertension
- Vasculitis?

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Not all MIS-C looks alike

**Presentations:
cardiogenic shock,
appendicitis, respiratory
failure, neurologic
decompensation**

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Multisystem Inflammatory Syndrome in Children (MIS-C)

Lab evidence of current or past infection with SARS-CoV-2



Fever, Myalgia
Conjunctivitis
Rash, Lymphadenopathy, Stomatitis, Extremity swelling with erythema
Skin peeling

Headache
Meningismus
Lethargy

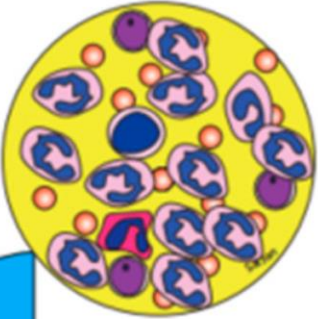
High ESR, CRP, ferritin, LDH, IL-6, Fibrinogen, Procalcitonin, CPK, D-dimers etc.,

Myocarditis, ↑Troponin, ↑pro-BNP
Coronary aneurysms, Hypotension
Hypoperfusion, Tachycardia

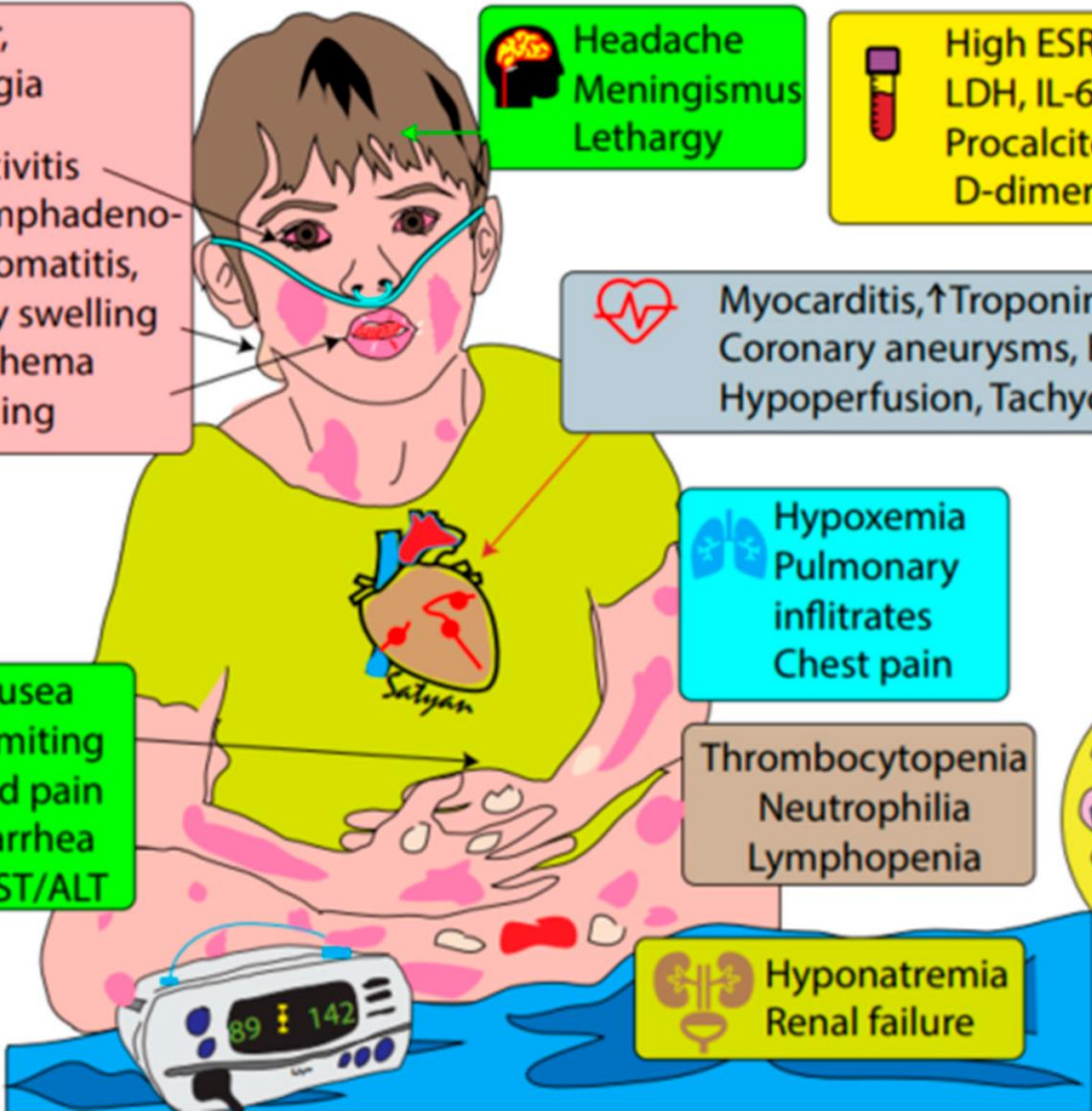
Nausea
Vomiting
Abd pain
Diarrhea
↑AST/ALT

Hypoxemia
Pulmonary infiltrates
Chest pain

Thrombocytopenia
Neutrophilia
Lymphopenia



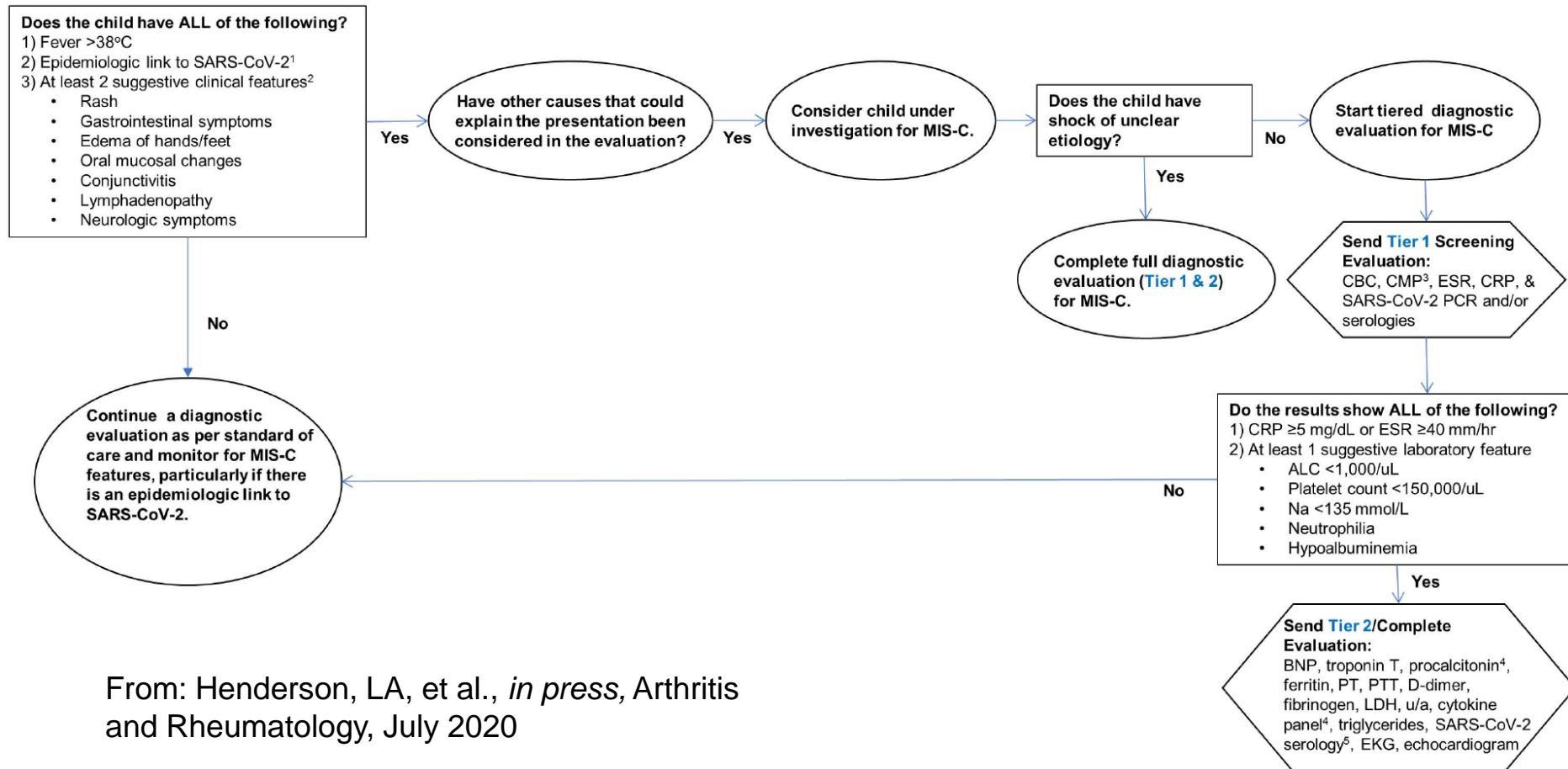
Hyponatremia
Renal failure



● Adapted from CDC

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Does my patient have MIS-C?



From: Henderson, LA, et al., *in press*, Arthritis and Rheumatology, July 2020

American College of Rheumatology Task Force

- Convened a group of experts: inclusive
 - 9 peds rheum, 2 cardiology, 1 ICU, 2 adult rheum
- Goals:
 - Provide clinical guidance, not guidelines
 - Target audience: pediatricians, pediatric specialists
- Based on scant evidence—creation of a “living” document that will be updated as evidence emerges
- Webinars and voting on strength of each recommendation
- 40 guidance statements

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Kawasaki disease vs. MIS-C

- $\frac{1}{4}$ to $\frac{1}{2}$ of MIS-C meet full criteria for KD
- Broad age distribution of MIS-C including many teens (KD usually younger <5)
- Presentation with shock unusual in KD
- Associated GI and neuro features less frequent in KD
- MIS-C—low platelets, higher CRP/ESR, lower lymph counts

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Therapy for MIS-C patients requiring ICU care

- General supportive care for organ dysfunction
- ECHO at presentation, repeat at 7-14 days
- Stepwise use of immunomodulatory treatment
 - IVIG or glucocorticoids as first line
 - May use in combination
 - Anakinra for patients who are refractory to above (or consideration of IL-6 blockade)
- Anti-platelet or anti-coagulation therapy
- Study is ongoing-BATS-Best Available Treatment Study

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