INTRODUCTION

The COVID-19 pandemic has caused significant disease across the globe but children seem to be much less affected than adults. Coincidentally with the first wave of the pandemic, a cluster of children with fever, hyperinflammation and shock were identified, and this was first described as paediatric inflammatory multisystem syndrome temporally associated with SARS-CoV-2 (PIMS-TS) by the Royal College of Paediatrics and Child Health. Patients with this novel condition were transferred to tertiary centres for management, increasing the pressure in these hospitals that were already extremely busy. There are multiple challenges related to the identification of patients presenting with PIMS-TS given that they mimic multiple other well-known paediatric conditions, like Kawasaki disease and toxic shock syndrome. Investigations and admission criteria include diagnostic uncertainty, access to investigations, the variable clinical phenotypes of PIMS-TS, high impact of complications and the requirement for specialist input. However, as our understanding of the condition has developed, more patients are now managed in a district general hospital (DGH), with virtual multidisciplinary team (MDT) involvement. Challenges include diagnostic uncertainty, access to investigations, the variable clinical phenotypes of PIMS-TS, high impact of complications and the requirement for specialist input.

Here we present a structured approach to management of children presenting with suspected PIMS-TS in a DGH.

DEFINITION

Children frequently present with a history of persistent fever, associated with gastrointestinal symptoms and/or mucocutaneous manifestations. Respiratory symptoms are infrequent (table 1). Many children seem relatively well at presentation but have disproportionately marked fever and/or unusual mucocutaneous features.

The median age is 9 years, but cases have been described as early as 3 months of age. There seems to be a male predominance, and most children...
have no underlying comorbidities.\textsuperscript{7} Black, Asian and minority ethnic (BAME) groups appear to be more affected, and SARS-CoV-2 exposure in the preceding 4–6 weeks was commonly found in these patients.\textsuperscript{4}

### Best practice

Results be positive or negative. SARS-CoV-2 infections associated with myocarditis (T wave inversion, pathological Q waves and ST changes).\textsuperscript{10} Additional investigations might be required depending on the severity of presentation (box 1). D-dimer results should be interpreted carefully, as there is little paediatric experience of its response to inflammation. All children with PIMS-TS should be invited to join the RECOVERY research study, other relevant studies recruiting locally (including DIAMONDS) and be

### INITIAL EVALUATION IN THE EMERGENCY DEPARTMENT

A thorough history and examination is vital to establish features of PIMS-TS, attempt to differentiate it from its clinical mimics and identify complications that may preclude the patient from being managed in the DGH.

Deciding which children to investigate can be challenging; children with early PIMS-TS are often deceptively well looking, but self-limiting febrile illness remains a more common presentation, and overinvestigation should be avoided. We suggest a pragmatic two-stage approach, supported by the national consensus pathway\textsuperscript{8} (figures 1 and 2).

There are reports of children who present appearing well and are discharged home, but re-present to hospital following rapid clinical deterioration.\textsuperscript{9} Therefore, robust safety-netting advice should be given to parents/carers.

Second-line investigations should include a chest radiograph and a 12-lead ECG, looking for signs of myocarditis (T wave inversion, pathological Q waves and ST changes).\textsuperscript{10} Additional investigations might be required depending on the severity of presentation (box 1). D-dimer results should be interpreted carefully, as there is little paediatric experience of its response to inflammation.

All children with PIMS-TS should be invited to join the RECOVERY research study, other relevant studies recruiting locally (including DIAMONDS) and be

### Figure 1

Decision-making chart of investigation.

### Table 1

<table>
<thead>
<tr>
<th>RCPCH definition of PIMS-TS</th>
<th>CDC definition of MIS-C</th>
</tr>
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<tbody>
<tr>
<td><strong>Child presenting with:</strong></td>
<td><strong>Individual of &lt;21 years with:</strong></td>
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<tr>
<td>Persistent fever, inflammation (neutrophilia, elevated CRP and lymphopaenia) and evidence of single or multiorgan dysfunction with additional features as indicated further.</td>
<td>At least 24 hours of subjective or objective fever &gt;38°C.</td>
</tr>
<tr>
<td><strong>Clinical features:</strong></td>
<td><strong>Severe illness needing admission.</strong></td>
</tr>
<tr>
<td>Fever &gt;38°C.</td>
<td>Two or more organ systems affected.</td>
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<tr>
<td>Abdominal symptoms: pain, diarrhoea and vomiting.</td>
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<tr>
<td>Rash/conjunctivitis.</td>
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<tr>
<td>Hypotension and shock.</td>
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<tr>
<td>Oxygen requirement.</td>
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<tr>
<td>Confusion.</td>
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<tr>
<td>Cough/respiratory symptoms.</td>
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<tr>
<td>Sore throat.</td>
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<td>Headache.</td>
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<tr>
<td>Mucous membrane changes.</td>
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<tr>
<td>Neck swelling.</td>
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<tr>
<td>Swollen hands and feet/ desquamation.</td>
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<tr>
<td>Syncope.</td>
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<tr>
<td><strong>Laboratory findings:</strong></td>
<td><strong>One or more of the following:</strong></td>
</tr>
<tr>
<td>Raised CRP.</td>
<td>Raised CRP, ESR, procalcitonin and IL-6.</td>
</tr>
<tr>
<td>Lymphopaenia/neutrophilia.</td>
<td>Lymphopaenia/neutrophilia.</td>
</tr>
<tr>
<td>Raised ferritin.</td>
<td>Low albumin.</td>
</tr>
<tr>
<td>Raised fibrinogen.</td>
<td>Raised ferritin.</td>
</tr>
<tr>
<td>Raised D-Dimer.</td>
<td>Raised fibrinogen.</td>
</tr>
<tr>
<td>Abnormal renal profile.</td>
<td>Raised D-Dimer.</td>
</tr>
<tr>
<td>Hyponatraemia.</td>
<td>Raised LDH.</td>
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<tr>
<td>Raised troponin and NT-proBNP.</td>
<td></td>
</tr>
<tr>
<td>This may include children fulfilling full or partial criteria for Kawasaki disease.\textsuperscript{*}</td>
<td>Some individuals may fulfil full or partial criteria for Kawasaki disease.\textsuperscript{*}</td>
</tr>
<tr>
<td>Exclusion of other microbial causes (bacterial sepsis, streptococcal/ staphylococcal shock syndromes, infections associated with myocarditis) but do not delay management while awaiting these results.</td>
<td>No alternative plausible diagnosis. Consider MIS-C in any paediatric death with evidence of SARS-CoV-2 infection.</td>
</tr>
</tbody>
</table>

**SARS-CoV-2 PCR test results may be positive or negative.**

| Positive for current or recent SARS-CoV-2 infection by RT-PCR, serology or antigen test or COVID-19 exposure within 4 weeks. | |

*Kawasaki disease criteria: 5 days of fever >4/5 clinical features: red rash, red conjunctival injection, red lips and mouth or cracked dry lips, red hands and feet and lymphadenopathy. CDC, Centers for Disease Control and Prevention; CRP, C reactive protein; ESR, erythrocyte sedimentation rate; IL-6, interleukin 6; LDH, lactic acid dehydrogenase; MIS-C, multisystem inflammatory syndrome in children; NT-proBNP, N-terminal pro-B type natriuretic peptide; RCPCH, Royal College of Paediatrics and Child Health; RT-PCR, reverse transcriptase PCR.

**Figure 2**

Investigations in children with suspected PIMS-TS. CRP, C reactive protein; ESR, erythrocyte sedimentation rate; LDH, lactic acid dehydrogenase; PIMS-TS, paediatric inflammatory multisystem syndrome temporally associated with SARS-CoV-2.

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Box 1  Additional investigations

- Abdominal ultrasound: if there is abdominal pain (±surgical input).
- Echocardiogram: when criteria for paediatric inflammatory multisystem syndrome temporally associated with SARS-CoV-2 are met.
- Save EDTA and serum for PCR and serological studies.
- Blood for SARS-CoV-2 serology.
- Blood and stool for SARS-CoV-2 PCR.
- Blood for pneumococcal, meningococcal, group A Strept and Staphylococcus aureus PCR.
- Antistreptolysin O titre and anti-DNase B.
- Epstein-Barr virus (EBV), cytomegalovirus (CMV), adenovirus and enterovirus PCR.
- Stool for virology.

<table>
<thead>
<tr>
<th>Table 2  Features of severe disease requiring transfer to tertiary unit</th>
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<tbody>
<tr>
<td>Prolonged capillary refill time</td>
</tr>
<tr>
<td>Hypotension/shock and/or tachycardia</td>
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<tr>
<td>Fluid resuscitation with 20 mL/kg fluid boluses or absent/paradoxical response</td>
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<tr>
<td>Respiratory failure</td>
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<tr>
<td>No response to IVIG/stereoids</td>
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CRITERIA FOR ADMISSION IN A DGH

Most children managed in the first wave of COVID-19 pandemic presented with severe disease and required transfer to a tertiary centre, with 44% needing PICU admission.

With increasing knowledge of the disease, the current management should be determined by the severity of the clinical picture and selected cases can be managed on a paediatric ward or high dependency unit of a DGH. These children may deteriorate rapidly, and close monitoring is vital. A febrile child with hyper-inflammation who is haemodynamically stable can be managed in a DGH, but should be discussed daily with their tertiary team (cardiology, rheumatology or infectious diseases, according to local pathways) to guide risk stratification and management decisions.

An echocardiogram should be performed in the first 24 hours of admission in stable children—if unavailable in the DGH, day transfer to a tertiary unit may be considered—and reviewed by a paediatric cardiologist. Daily bloods including inflammatory markers, clotting profile and troponin/NT-proBNP should be done.

CRITERIA FOR TRANSFER TO A TERTIARY CENTRE

All children who present severely ill, with shock and multiorgan failure, should be retrieved to a PICU. Patients who present with concerning features should also be discussed with tertiary centres for potential transfer (table 2). Children with PIMS-TS who develop shock are challenging to manage in a DGH. They can develop profound hypotension and respond unpredictably to fluids. Early PICU advice and retrieval is strongly recommended.

Cardiovascular instability is the most frequent factor in acute decompensation. Many children have fluid-unresponsive shock or develop signs of cardiac failure with fluid. Consider starting with 5–10 mL/kg bolus and if subsequent boluses are needed, and there are concerns of reduced cardiac function, give 5 mL/kg aliquots to allow more frequent assessments of the response. Inotropes/vasopressors are often needed, and the local team should be prepared to start these before the arrival of the retrieval team. The choice should be guided by clinical condition and local pathways. In some cases mechanical ventilation is required, mostly for cardiovascular support rather than respiratory failure.

PIMS-TS also has several effects on the heart caused by myocarditis. The wide range of manifestations include pericardial effusion, mild valvar regurgitations, coronary artery dilatation and myocarditis with left ventricular dysfunction. Regular echocardiograms are needed to assess severity; this might be challenging to obtain in a DGH. Therefore, children with cardiac involvement should be managed in a tertiary centre.

DIFFERENTIAL DIAGNOSIS

This novel condition manifests as a hyperinflammatory response with multiorgan involvement, with overlapping features of Kawasaki disease (KD), staphylococcal and streptococcal toxic shock syndrome (TSS) (figure 3) and macrophage activation syndrome/haemophagocytic lymphohistiocytosis. Other differentials include invasive group A streptococcal disease, invasive staphylococcal disease, other bacterial sepsis and viral infections (measles, EBV, adenovirus, parvovirus and enterovirus) and acute abdomen such as appendicitis.

HOW IS PIMS-TS DIFFERENT FROM OTHER CONDITIONS?

PIMS-TS patients are different from KD and TSS patients, as summarised in figure 4.
However, three distinctive classes of patients can be identified based on the clinical phenotype: PIMS-TS only, PIMS-TS/KD and PIMS-TS/TSS.

Children with PIMS-TS solely have the least severe presentation, with median age of 7.8 years. Within the group with PIMS-TS/KD, children are younger (median age 4.5 years) and they have the highest proportion of skin involvement, but criteria for KD (complete or incomplete) are not met in about 60% of the cases. The incidence of coronary aneurysms seems to be much higher in KD than in PIMS-TS. Furthermore, overlapping features between these syndromes suggest that they may share similar pathophysiology and likely explains why these patients respond to similar therapies.

**MANAGEMENT**

**Initial stabilisation**

On initial assessment of a child who presents to the hospital with persistent fever, inflammation and evidence of single or multiorgan dysfunction and/or Kawasaki-like features, appropriate personal protective equipment (PPE) should be worn and resuscitation and supportive management started.

**Immunomodulation**

In stable patients with suspected PIMS-TS, if consent is not obtained for the RECOVERY trial, intravenous immunoglobulin (IVIG) should be given unless advised otherwise by the tertiary hospital. Second-line modulation is intravenous methylprednisolone, to be considered in patients remaining unwell 24 hours after IVIG.

**Figure 3** Differential diagnosis of PIMS-TS. PIMS-TS, paediatric inflammatory multisystem syndrome temporally associated with SARS-CoV-2.

**Figure 4** Unique characteristics of PIMS-TS. GI, gastrointestinal; CRP, C reactive protein; CVS, cardiovascular; KDSS, Kawasaki Disease shock syndrome; PIMS-TS, paediatric inflammatory multisystem syndrome temporally associated with SARS-CoV-2.
particularly with ongoing fever. According to the Best Available Treatment Study, no difference in outcomes was found between treatment with glucocorticoids or IVIG as single agents or between the single-agent and dual-agent primary treatments.16

If the child’s condition deteriorates or they do not respond to IVIG or methylprednisolone, then biologicals should be considered (tocilizumab, infliximab and anakinra) and the child should be transferred to a tertiary centre.15

We recommend the use of methylprednisolone as first line in haemodynamically unstable patients for two reasons: first, it contributes to maintaining blood pressure and second, there is a risk of volume overload with IVIG therapy, which could lead to acute decompensation.

**DISCHARGE CRITERIA AND FOLLOW-UP**
Children admitted to the DGH with PIMS-TS can be considered for discharge once they remain afebrile for over 24 hours, have stable cardiac function and decreasing inflammatory markers. Follow-up should be arranged for 1–2 weeks after discharge and at 6 weeks. An echocardiogram should be performed as part of follow-up, since coronary artery aneurysms may develop even after mild cases.8

**Additional support**
Play therapists should be involved in management to reduce anxiety/fear associated with prolonged hospital stays, painful procedures and staff wearing PPE.
Best practice

Test your knowledge

1. Which of the following are true regarding epidemiology of paediatric inflammatory multisystem syndrome temporally associated with SARS-CoV-2 (PIMS-TS)? You may select more than one answer.
   A. The median age at presentation is 12 years.
   B. There is a slight female predominance.
   C. Most children have an underlying comorbidity.
   D. Black, Asian and minority ethnic (BAME) groups are more commonly affected.
   E. Cases have been identified in children as young as 3 months of age.

2. Which of the following is true regarding investigations for suspected PIMS-TS?
   A. Abdominal ultrasound should be performed in all patients with suspected PIMS-TS.
   B. Stable children may be suitable for day transfer to a tertiary unit for echocardiogram if this is not available in the district general hospital (DGH).
   C. First-line investigation should include ferritin, D-dimer and troponin.
   D. Negative SARS-CoV-2 antibody testing excludes a diagnosis of PIMS-TS.
   E. Second-line investigations should only be performed if initial investigations show significant neutrophilia, lymphopaenia, thrombocytopaenia and/or CRP rise.

3. Which of the following findings are much more commonly seen in Kawasaki disease than PIMS-TS? You may select more than one answer.
   A. Abdominal pain.
   B. Coronary artery aneurysm.
   C. Mucosal changes.
   D. Raised C reactive protein (CRP).
   E. Signs of cardiovascular shock.

4. A 10-year-old boy with no significant comorbidities presents to the emergency department in a DGH with a 4-day history of fever, lethargy, rash, sore throat and abdominal pain. There is no clear focus for infection, but he is slightly tachycardic with his fever. He is otherwise haemodynamically stable. Intravenous access is obtained and initial investigations show raised CRP, low lymphocytes and raised neutrophils. Which of the following are true regarding his management?
   A. He should be discussed with a tertiary centre before commencing any further treatment.
   B. Intravenous fluid should be given in 20 mL/kg aliquots in view of his tachycardia.
   C. If he remains haemodynamically stable, ceftriaxone alone should be commenced, along with human intravenous immunoglobulin, unless consent is obtained for the RECOVERY trial.
   D. He should receive low dose aspirin and proton pump inhibitor.
   E. If his chest X-ray, ECG and troponin are normal, there is no indication for echocardiogram.

Test your knowledge

5. Which of the following are required when considering children for discharge after treatment for suspected PIMS-TS?
   A. Afebrile >24 hours.
   B. Falling inflammatory markers.
   C. Stable cardiac function.
   D. Plans for outpatient follow-up including echocardiogram.
   E. All of the above.

Answers to the quiz are at the end of the references.

FUTURE DIRECTIONS
A growing understanding of PIMS-TS and its management means that more patients can be managed effectively in the DGH. This reduces the need for transfer and pressure on tertiary centres and allows families to stay closer to home. Virtual MDT meetings and well-developed protocols have fostered a safe, collaborative relationship between the DGH and tertiary units, resulting in better patient care. We hope that the lessons learnt through this process can be applied to other conditions in the future.

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REFERENCES


Answers to the multiple-choice questions
1. D and E.
2. B.
3. B and C.
4. C and D.
5. E.