Covid-19 and acute kidney injury in hospital: summary of NICE guidelines

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What you need to know

- Acute kidney injury (AKI) may be common in patients with covid-19 and is associated with an increased risk of dying
- AKI associated with covid-19 may be caused by volume depletion, multi-organ failure, viral infection leading directly to kidney tubular injury, thrombotic vascular processes, glomerulonephritis, or rhabdomyolysis
- Maintaining optimal fluid status (euvolaemia) is critical in reducing the incidence of AKI
- Regular assessments of fluid status and fluid management plans are necessary, and in those who need intravenous fluids the choice of replacement fluid should be based on patients’ biochemistry and fluid status
- An increased risk of coagulopathy may cause problems with clotting of the extracorporeal circuit during renal replacement therapy

Acute kidney injury (AKI), a sudden reduction in kidney function, is seen in some people with covid-19 infection. A subset of patients develop severe AKI and require renal replacement therapy (RRT). As in many settings, the development of AKI is associated with an increased risk of mortality. Although our understanding is incomplete, a picture is emerging from case reports and autopsy series of covid-19 specific causes of AKI. Intrinsic renal pathology including thrombotic vascular processes, viral mediated tubular cell injury, and glomerulonephritis have been reported, as well as AKI resulting from extrinsic factors such as fluid depletion, multi-organ failure, and rhabdomyolysis. Anecdotal reports have emerged of proximal tubular injury with Fanconi syndrome that manifests as hypokalaemia, hypophosphataemia, normal anion gap metabolic acidosis, and hypovolaemia from salt wasting. Importantly, AKI can occur at all stages of covid-19 infection, so clinical vigilance and consideration of risk factors for AKI alongside early detection and diagnosis are essential components of general supportive care. Fluid management is central to this.

This article summarises key points from the National Institute for Health and Care Excellence (NICE) covid-19 rapid guideline on AKI in hospital.

Recommendations

Communicating with patients

- Communicate effectively with patients, their families, and carers, and support their mental wellbeing to help alleviate any anxiety they may have about covid-19. Signpost to charities and UK government guidance on the mental health and wellbeing aspects of covid-19.

Minimising risk for patients and healthcare workers

- All healthcare workers involved in receiving, assessing, and caring for patients who have known or suspected covid-19 should follow UK government guidance for infection prevention and control.
- If covid-19 is later diagnosed in a patient not isolated from admission or presentation, follow UK government guidance on management of exposed healthcare workers and patients in hospital settings.

Planning treatment and care

- Discuss the risks, benefits, and likely outcomes of treatment options with patients with covid-19, and their families and carers. This will help them make informed decisions about

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their treatment goals and wishes, including treatment escalation plans where appropriate.

- Find out if patients have advance care plans or advance decisions to refuse treatment, including “do not attempt cardiopulmonary resuscitation” decisions, and take account of these in planning care.
- Monitor patients for the development or progression of chronic kidney disease (CKD) after AKI. Guidance on care after hospital discharge was produced jointly by Think Kidneys and the Royal College of General Practitioners is designed to support safer transitions of care and post-discharge monitoring, and is of relevance to both hospital and general practice teams.\(^1\)

**Assessing for AKI in patients with suspected or confirmed covid-19**

Be aware that, in patients with covid-19, AKI

- may be common, but prevalence is uncertain and depends on clinical setting; the Intensive Care National Audit and Research Centre’s report on covid-19 in critical care reported that 31% of patients on ventilators and 4% not on ventilators needed renal replacement therapy for AKI\(^2\)
- is associated with an increased risk of dying\(^1,2\)
- can develop at any time before or during hospital admission
- causes may include volume depletion (hypovolaemia), haemodynamic changes, viral infection leading directly to kidney tubular injury, thrombotic vascular processes, glomerular pathology, or rhabdomyolysis\(^3\)
- may be associated with haematuria, proteinuria, and abnormal serum electrolyte levels (both increased and decreased serum sodium and potassium).\(^4\)

Be aware that in patients with covid-19

- maintaining optimal fluid status (euvoaemia) is critical in reducing the incidence of AKI, but this can be hard to achieve
- treatments being used to manage covid-19 may increase the risk of AKI—for example, diuretics if they have caused volume depletion (hypovolaemia)
- fever and increased respiratory rate increase insensible fluid loss
- dehydration (often needing correction with intravenous fluids) is common on admission to hospital and may also develop later
- risk of coagulopathy is increased.

On hospital admission or transfer, assess for AKI in all patients.

Record

- medical history and comorbidities, including factors that further increase the risk of AKI (such as CKD, heart failure, liver disease, diabetes, history of AKI, age 65 or over)
- fluid status by clinical examination (for example, peripheral perfusion, capillary refill, pulse rate, blood pressure, postural hypotension, jugular venous pressure, or pulmonary or peripheral oedema)
- fluid status by fluid balance (fluid intake, urine output, and weight)
- full blood count
- serum urea, creatinine, and electrolytes (sodium, potassium, bicarbonate).

Review the use of medicines that can cause or worsen AKI and stop these unless essential.

- Ask a pharmacist for advice about optimising the choice and dosage of medicines, including anticoagulants for treatment or prophylaxis. More detailed information is available in the Think Kidneys guidelines for medicines optimisation in patients with AKI.\(^5\)

Continue to assess for AKI. Record and monitor fluid status by clinical examination and fluid balance daily. Measure serum urea, creatinine, and electrolytes (sodium, potassium, bicarbonate) at least every 48 hours or more often if clinically indicated (eg, in those at increased risk of AKI, in those who have sustained AKI, and those with electrolyte abnormalities). Use an early warning score for patients whose clinical condition is deteriorating or who have suspected sepsis:

- NEWS2 has been endorsed by NHS England.
- When using NEWS2 be aware of the Royal College of Physicians’ warning that any increase in oxygen requirements should be escalated for clinical review and increased observations.\(^6\)

**Detecting and investigating AKI in patients with suspected or confirmed covid-19**

Detect AKI using NHS England’s AKI algorithm\(^7\) or any of the following criteria:

- an increase in serum creatinine of ≥26 μmol/L in 48 hours
- an increase of ≥50% in serum creatinine, known or presumed to have occurred in the past seven days
- a fall in urine output to ≤0.5 mL/kg/hour for more than six hours.

Do urinalysis for blood, protein, and glucose to help identify the cause of AKI. Record the results and take action if these are abnormal (including referral if needed; see section below on referral in patients with suspected or confirmed covid-19).

Perform imaging if urinary tract obstruction is suspected.

**Managing fluid status in patients with suspected or confirmed covid-19**

- Aim to achieve and maintain optimal fluid status (euvoaemia) in all patients.
- If there is volume depletion (hypovolaemia) and fluid needs cannot be met orally or enterally, give patients intravenous fluids as part of a protocol to restore and maintain optimal fluid status (euvoaemia).
- Ensure patients have an intravenous fluid management plan that is reviewed daily.
- Base choice of fluids on biochemistry results and fluid status. The composition of commonly used fluids is summarised in table 1.
- Do not routinely offer loop diuretics to treat AKI but consider them for treating fluid overload.

**Managing hyperkalaemia in patients with suspected or confirmed covid-19**

- Be aware of the risk of hyperkalaemia and manage according to local protocols.
- The potassium binders patiromer and sodium zirconium cyclosilicate can be used alongside standard care for the emergency management of acute life threatening
hyperkalaemia (these agents have been approved by NICE for this indication).19,20

**Referral in patients with suspected or confirmed covid-19**

Refer patients with AKI for further specialist advice if:
- There is diagnostic uncertainty about the cause of AKI, which may need further tests or imaging.
- They have abnormal urinalysis results, which may be a sign of COVID-19-induced kidney damage or other intrinsic renal disease.
- Fluid management needs are complex.
- AKI is worsening despite initial management or has not resolved after 48 hours.
- The patient has usual indications for renal replacement therapy, particularly if there is no urine output, such as:
  - Life-threatening hyperkalaemia.
  - Refractory fluid overload.
  - Severe metabolic acidosis.

**Renal replacement therapy in patients with suspected or confirmed covid-19**

The scope of the guideline did not include a detailed review of the technical aspects of provision of renal replacement therapy (RRT) in COVID-19. Resources were signposted as follows:

- NHS England has produced a clinical guide on renal replacement therapy options in critical care during the coronavirus pandemic for options for patients with usual indications for RRT based on local availability, equipment, supplies, staffing, and local expertise.21
- The Renal Association has collated a set of COVID-19 resources, which include protocols for RRT.22
- Be aware of the anecdotal reports of RRT circuit clotting because of the increased risk of coagulopathy in patients with COVID-19.
- No evidence was found on how best to provide anticoagulation during RRT in patients with COVID-19.

**Areas of uncertainty**

Information regarding renal involvement in COVID-19 is extremely limited in several areas, and further evidence is required. Some of the most pressing questions include:

- What is the incidence of AKI in hospitalised patients with COVID-19, both in and outside of the intensive care unit?
- What, if any, are the typical clinical, laboratory, and urinary features that characterise AKI in the setting of COVID-19?
- What are the different histological patterns of renal involvement in COVID-19 and how do these relate to clinical presentation?
- What are the long-term effects of COVID-19 on renal function, including the proportion of survivors who require ongoing renal replacement therapy resulting from end-stage kidney disease?

**Guidelines into practice**

- Can you identify patients with COVID-19 who are at particular risk of sustaining AKI?
- Do you know which patients with COVID-19 associated AKI should be referred for specialist advice, and do you know your local referral pathway?
- How should patients who have sustained COVID-19 associated AKI be followed up in primary care, and do you know where to find RCPGP guidance on AKI care after hospital discharge?


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## Table

### Table 1: Composition of commonly used fluids, adapted from NICE Guideline (CG174)

<table>
<thead>
<tr>
<th>Content</th>
<th>Plasma Sodium chloride 0.9%</th>
<th>Sodium chloride 0.18%/4% glucose</th>
<th>Sodium chloride 0.45%/4% glucose</th>
<th>5% glucose Hartmann’s Ringer’s</th>
<th>Ringer’s acetate</th>
<th>Alternative balanced solutions for resuscitation**</th>
<th>Alternative balanced solutions for maintenance**</th>
</tr>
</thead>
<tbody>
<tr>
<td>Na⁺ (mmol/L)</td>
<td>135-145</td>
<td>154</td>
<td>31</td>
<td>77</td>
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<td>131</td>
<td>130</td>
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<tr>
<td>Cl⁻ (mmol/L)</td>
<td>95-105</td>
<td>154</td>
<td>31</td>
<td>77</td>
<td>0</td>
<td>111</td>
<td>109</td>
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<tr>
<td>Na⁺:Cl⁻ ratio</td>
<td>1.28-1.45:1</td>
<td>1:1</td>
<td>1:1</td>
<td>1:1</td>
<td>-</td>
<td>1.18:1</td>
<td>1.19:1</td>
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<tr>
<td>K⁺ (mmol/L)</td>
<td>3.5-5.3</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>5</td>
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<tr>
<td>HCO₃⁻ (mmol/L)</td>
<td>24-32</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>29 (lactate)</td>
<td>28 (lactate)</td>
<td>27 (acetate)</td>
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<tr>
<td>Ca²⁺ (mmol/L)</td>
<td>2.2-2.6</td>
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<td>0</td>
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<td>2</td>
<td>1.4</td>
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<td>Mg²⁺ (mmol/L)</td>
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<td>1</td>
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<td>Glucose (mmol/L)</td>
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<td>222</td>
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<td>pH</td>
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<td>4.5</td>
<td>3.5-5.5</td>
<td>5.7</td>
<td>6-7.5</td>
<td>6-8</td>
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<td>Osmolarity (mOs/m/L)</td>
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<td>308</td>
<td>284</td>
<td>278</td>
<td>278</td>
<td>273</td>
<td>276</td>
</tr>
</tbody>
</table>

* These solutions are available with no potassium or differing quantities of potassium already added

** Alternative balanced solutions are available commercially under different brand names and composition may vary by preparation