Secondary sclerosing cholangitis in critically ill patients: a rare disease precipitated by severe SARS-CoV-2 infection

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SUMMARY
A previously well 59-year-old man required a prolonged intensive care unit stay due to severe COVID-19 symptoms. During the admission, he developed a cytokine storm, also known as secondary haemophagocytic lymphohistocytosis, and multiorgan failure. Despite recovering from his other organ failures, his liver function continued to deteriorate. Magnetic resonance cholangiopancreatography and subsequent endoscopic retrograde cholangiopancreatography revealed extensive intrahepatic bile duct dilatation with 'beading' but common bile duct sparing. Given the patient had no primary liver disease prior to admission, we considered secondary causes of cholestatic liver injury; this led us to an unusual diagnosis of secondary sclerosing cholangitis in critically ill patients. This case demonstrates a rare disease that has developed specifically in the context of SARS-CoV-2 infection. A review of current literature and the underlying pathophysiology for this rare disease are discussed, particularly in relation to COVID-19.

BACKGROUND
Secondary sclerosing cholangitis (SSC) is diagnosed when a cause for progressive cholestatic liver injury is identified in patients who have no prior history of hepatobiliary disease. Several secondary causes have been identified over recent years, including ischaemia and iatrogenic, yet the SSC in critically ill patients (SSC-CIP) subgroup is particularly rare and was first described in 2001 by Leonhardt et al.

The underlying pathophysiology in patients with SSC-CIP is thought to be a combination of bile duct ischaemia and changes to the composition of bile. This results in cholangioocyte necrosis and formation of bile casts that obstruct the biliary system, giving rise to recurrent cholangitis.

The most important factor distinguishing SSC-CIP from other hepatobiliary diseases related to critically ill patients is the persistent cholestasis despite clinical recovery of other organ injuries. This progressive cholestasis is reflective of irreversible cellular damage, and hence a relatively poor prognosis often with rapid deterioration to biliary cirrhosis, requiring transplantation as an only means of cure.

SSC-CIP is not widely appreciated by physicians and intensivists, such that it may go unrecognised in some patients. This case therefore provides an excellent learning opportunity to inform readers of this rare disease, particularly in the context of COVID-19, with the aim of promoting earlier disease recognition to allow timely investigation and optimal management strategies, including referral for transplant assessment.

Here a rare case of SSC-CIP is reported after a prolonged intensive care admission due to SARS-CoV-2 infection.

CASE PRESENTATION
A previously fit and well 59-year-old man was admitted to the intensive care unit (ICU) with a 2-week history of cough, fever and breathlessness. Admission blood profile revealed lymphopenia, with a d-dimer of >20000 μg/L and ferritin of 3991 μg/L. He was intubated and ventilated for severe hypoxia on day 1 and was subsequently confirmed SARS-CoV-2 positive via oropharyngeal swab. Shortly after admission, the patient developed a cytokine storm, known as secondary haemophagocytic lymphohistocytosis (sHLH) and multiorgan failure. He was also treated with levofloxacin for presumed superimposed bacterial pneumonia. During his ICU admission, the patient was also treated for staphylococcus septicemia and cellulitis with vancomycin and co-trimoxazole and later developed Candida albicans infection from the central line, bile and urine cultures, which was managed with a course of caspofungin. He intermittently required inotrope support and developed necrotic toes on his left foot. The patient was slow to wake up from his sedation and had a prolonged respiratory wean, leading to tracheostomy insertion 3 weeks into his admission, with a successful decanulation 1 month later.

While in ICU, it was noted that the patient’s liver function tests (LFTs) were becoming deranged with a cholestatic pattern, having been normal on admission since the date of discharge.

INVESTIGATIONS
Imaging while on ICU included two abdominal ultrasound scans, which did not reveal any parenchymal or biliary tree abnormality. However, a magnetic resonance cholangiopancreatography (MRCP) undertaken when the patient was discharged from ICU revealed a few hypointense filling defects within the common bile duct (CBD) suspicious of choledocholithiasis. The intrahepatic bile ducts were also dilated and demonstrated some
beading (figure 2). A subsequent endoscopic retrograde cholangiopancreatography (ERCP) revealed ‘bizarre appearances’ with a large column of sludge within the whole CBD and a sclerosing cholangitis type picture within the intrahepatic ducts (figure 3). As part of a non-invasive liver screen, serology revealed negative Anti-Neutrophilic Cytoplasmic Autoantibody (ANCA), IgG-4 and liver autoantibodies.

DIFFERENTIAL DIAGNOSIS

When this man was initially reviewed on ICU, the deteriorating LFTs were thought to be caused by a combination of pathophysiological factors accumulating in multiorgan failure, driven by COVID-19 and superimposed bacterial septicaemia. At that time, given the normal appearance on ultrasound scanning, a ‘watch and wait’ approach was taken on the assumption the LFTs would normalise as the patient improved clinically. However, despite other organ systems recovering, the LFTs continued to deteriorate. Differentials at this point were vast, however, primary liver disease, such as autoimmune hepatitis or primary sclerosing cholangitis (PSC), was low on the list of differentials given the patient had no known hepatobiliary disease prior to admission. Hence, we considered secondary causes of cholestatic liver injury in the context of critically unwell adults; drug-induced and whether or not referral to a regional transplant centre will be required.

TREATMENT

The patient underwent two ERCP procedures; on the first occasion, endoscopic sphincterotomy and balloon trawl of sludge were performed. His bilirubin peaked at 241 μmol/L just before this procedure and slowly fell during subsequent weeks. A trial of ursodeoxycholic acid was also initiated after the first procedure. A further ERCP and balloon trawl of sludge with bile taken for culture were performed 1 month later. At the time of write-up, the patient is awaiting liver biopsy to further evaluate the degree of hepatocyte and sinusoidal injury.

OUTCOME AND FOLLOW-UP

The patient, who’s admission was precipitated by SARS-CoV-2 infection, has since been discharged from the secondary care setting. Liver biopsy will further inform as to the degree of resulting parenchymal and portal tract architectural disturbance, and whether or not referral to a regional transplant centre will be required.

DISCUSSION

Unlike PSC, which is an idiopathic disease process,6 SSC is diagnosed when a cause for the progressive cholestatic injury is identified in patients with no prior history of hepatobiliary disease.1 Many secondary causes have been identified over recent years inclusive of ischaemia, drug-induced and auto-immune IgG-4,7 however, the SSC-CIP subgroup is a particularly rare disease that was first reported in 2001 by Scheppach et al.8 It has an estimated prevalence of 1 in 2000 ICU admissions, but only 250 cases have so far been reported worldwide.1 The patients documented with SSC-CIP have all been treated in an ICU setting for multiple reasons including that of severe infection, trauma, burns and postcardiothoracic surgery.8 Typically, in such patients, the cholestatic pattern of deranged LFTs persists beyond the recovery of the disease that initially precipitated ICU admission.1

The underlying pathophysiology in patients with SSC-CIP is not completely understood but is generally thought to be due to a combination of bile duct ischaemia (ischaemic cholangiopathy) and changes to the composition of bile (toxic bile), which together result in cholangiocyte necrosis and cast formation.2 This relative bile duct obstruction can precipitate infection, thus causing further disease progression to irreversible intrahepatic bile duct destruction and subsequent secondary biliary cirrhosis.2 Both ischaemic cholangiopathy and toxic bile formation can develop for multiple reasons relating to critical illness and management within an ICU setting.

Figure 1 Cumulative figures of serum bilirubin, alanine transaminase (ALT) and alkaline phosphatase (ALP) concentrations throughout the patient’s secondary care admission. The green vertical line demonstrates discharge from intensive care unit (ICU), and the red vertical line demonstrates discharge from secondary care.

Figure 2 Magnetic resonance cholangiopancreatography image showing ‘beading’ effect of intrahepatic bile ducts.

Figure 3 Cholangiogram taken during endoscopic retrograde cholangiopancreatography revealing a sclerosing cholangitis type picture within the intrahepatic ducts.
Ischaemic cholangiopathy

Intrahepatic biliary epithelium is particularly susceptible to ischaemia because anatomically it only has a singular blood supply via the hepatic arteries, unlike the CBD and hepatocytes that have a dual blood supply. Bile duct ischaemia is thought to be important for several reasons within a critical care setting; macrocirculatory haemodynamic instability with reduced mean arterial pressures requiring vasopressor support occurs in up to a third of patients admitted to ICU; this figure ranges between 60% and 100% in those patients developing SSC-CIP. While vasopressors help to increase systemic blood pressure, they often have a negative effect on hepatosplanchnic blood supply; for example, norepinephrine, which is commonly used in ICU to aid haemodynamic stability, has a vasoconstrictive property that reduces splanchnic blood flow, thus precipitating relative ischaemia.

Additionally, microcirculatory compromise within the peribiliary vascular plexus is a significant feature of SSC-CIP; Deltenre and Valla demonstrated that the degree of ischaemic cholangiopathy was inversely proportional to the calibre of the supplying occluded artery. According to Leonhardt et al., reasons for microcirculatory blood flow disturbance when a patient is critically unwell include increased blood viscosity and hypercoagulable states. This is particularly relevant in the context of severe SARS-CoV-2 infection in which a number of alterations in prothrombotic factors have been demonstrated, resulting in a hypercoagulable state. Hyperviscosity, which promotes a hypercoagulable state, has also been reported by Maier et al. in patients with COVID-19 who require critical care intervention. Hypercoagulable states in the context of SARS-CoV-2 infection, known as COVID-19-associated coagulopathy, have demonstrated a higher risk of both venous and arterial thromboembolism. Given that the predominant clinical finding in patients with COVID-19 is thrombosis, rather than bleeding, helps distinguish this pathological process from a disseminated intravascular coagulation-like state.

Arterial thrombosis has been documented in patients infected with SARS-CoV-2, whereby D-dimer levels have been recorded in excess of 9000 µg/L. This is reflective of the patient described in our case study whose D-dimer exceeded 20000 µg/L on admission. Formation of microvascular thrombi has also been demonstrated in post-mortem studies of those who have died with severe COVID-19 infection, the underlying cause of which is thought to be linked to hypercoagulopathy, direct endothelial damage and hyperviscosity states.

Mechanical ventilation with high positive end-expiratory pressures (PEEP) greater than 10 cm H₂O has also been shown to contribute to microcirculatory ischaemia within the hepatosplanchnic vascular plexus. Additionally, excessive use of prone positioning of mechanically ventilated patients has been linked to the development of SSC-CIP. Both these factors are particularly relevant in the context of severe SARS-CoV-2 infection, whereby mechanical ventilation with high PEEP for prolonged periods due to the challenges of weaning plus the use of proning in such patients for up to 16 hours per day are relatively common.

Overall, disturbances in the arterial supply of the peribiliary vascular plexus lead to cholangiocyte necrosis with subsequent formation of biliary casts and inflammation of intrahepatic bile ducts. This results in cholestasis and recurrent biliary infection, which risks further biliary obstruction and cirrhosis through ongoing scarring and inflammation.

Toxic bile

Toxic bile is also implicated as a causative factor of SSC-CIP. Usually defence mechanisms rely on hepatobiliary transporters to protect cholangiocytes from the toxic bile salts. Ischaemia predisposes to failure of such transporter systems which subsequently contributes to cholestasis and cell necrosis. An additional defence mechanism that is adversely affected by ischaemia is that of bicarbonate secretion via the chloride–bicarbonate exchanger 2 (AE2). Normally, this process promotes an alkaline pH on the apical surface of cholangiocytes and thus prevents permeation of bile acids. In addition, proinflammatory cytokines have been shown to inhibit the activity of AE2. This is important in the context of severe SARS-CoV-2 infection due to its association with a syndrome of uncontrolled immune activation leading to a cytokine storm, also known as HLH. Indeed Leonhardt et al. found that all 16 patients in their study had features of a systemic hyperinflammatory syndrome prior to their diagnosis of SSC-CIP. This further suggests that a heightened systemic inflammatory response through the release of proinflammatory cytokines adds to the development of toxic bile, hence contributing to cholangiocyte necrosis, and as such has an important association with SARS-CoV-2 infection.

Diagnosing SSC-CIP is challenging for several reasons, particularly given the disease is not well recognised and the underlying pathophysiological mechanisms are not completely understood. Also, patients with SSC-CIP are largely asymptomatic initially, presenting only with a cholestatic pattern of deranged LFTs, for which there are many more common differentials in patients who are critically unwell. Here, the differentials that one must consider include sepsis cholestasis, choledocholithiasis, drug-induced liver injury or adverse effects of intravenous nutrition.

While ultrasound imaging can be undertaken at the bedside, it is operator dependent and cannot always be relied on in the diagnosis of intrahepatic biliary diseases; while some authors report visible changes within the hepatobiliary complex in accordance with sclerosing cholangitis, others have reported unremarkable ultrasound imaging in keeping with the case discussed. MRCP or ERCP are more likely to aid the diagnosis of SSC-CIP, but these investigations are often delayed due to patients not being sufficiently stable to be moved out of the critical care environment. Early MRCP findings comprise of intrahepatic biliary filling defects due to accumulation of biliary casts. This subsequently progresses to diffuse biliary stricturing and dilatations with the classic beaded appearance. The main hallmark however of SSC-CIP that differentiates it from other potential diagnoses is that the extrabiliary hepatic system is typically spared. MRCP and/or ERCP remain the gold standard in imaging for patients with suspected SSC-CIP.

SSC-CIP carries with it a relatively poor prognosis; mortality rates in such patients are as high as 50% during an ICU admission, and adverse prognostic factors include associated renal failure, higher model for end-stage liver disease (MELD) scores and rapid deterioration to liver cirrhosis. A study by Lin et al. revealed that 60% of patients with SSC-CIP survived ICU admission; of these, 40% developed stable biliary cirrhosis and the remaining 20% progressed to transplantation. Without transplant, median survival in such patients is 12–44 months, compared with 89 months for patients with PSC.

Endoscopic procedures to aid biliary drainage such as sphincterotomy, balloon dilatation and stenting may bring about a transient improvement in clinical and biochemical status. As such, sphincterotomy and balloon trawl of sludge may have helped in our patient. Ursodeoxycholic acid has also been used to

improve hyperbilirubinaemia, but its effect is somewhat limited in SSC-CIP cases. Importantly, however, these interventions do not affect prognosis, with orthoptic liver transplantation being the only cure once biliary cirrhosis is established. Appropriateness of allocation of liver transplantation is primarily based on model for end stage liver disease (MELD) or United Kingdom model for end-stage liver disease (UKELD) scoring, with 75% of patients with SSC-CIP referred within 12 months of initial diagnosis. Survival rates post-transplant are reassuring at 85% after 3 years.

Learning points

► Consider secondary sclerosing cholangitis in critically ill patients (SSC-CIP) as a differential diagnosis for those critically unwell patients who have a persistent cholestasis despite clinical recovery of other organ failures.

► Ultrasound imaging cannot be relied on in patients with SSC-CIP, and therefore patients require further imaging with MRCP and/or endoscopic retrograde cholangiopancreatography as gold standard.

► The main hallmark of SSC-CIP that differentiates it from other potential diagnoses is that the extrahepatic biliary system is typically spared.

► Prognosis is relatively poor with rapid progression to biliary cirrhosis. Endoscopic procedures for biliary drainage and ursodeoxycholic acid do not alter the prognosis.

► The only cure for SSC-CIP is liver transplantation, and referral should be guided primarily by the model for end-stage liver disease (MELD) or United Kingdom model for end-stage liver disease (UKELD) scoring.

REFERENCES


