

## Care of patients with liver disease during the COVID-19 pandemic:

### EASL-ESCMID Position Paper

Tobias Boettler<sup>1</sup>, Philip N Newsome<sup>2,3</sup>, Mario U. Mondelli<sup>4</sup>, Mojca Maticic<sup>5,6</sup>, Elisa Cordero<sup>7</sup>,  
Markus Cornberg<sup>8,9</sup>, Thomas Berg<sup>10,\*</sup>

*<sup>1</sup>Department of Medicine II, Medical Center – University of Freiburg, Faculty of Medicine, University of Freiburg, Freiburg, Germany; <sup>2</sup>National Institute for Health Research, Birmingham Biomedical Research Centre at University Hospitals Birmingham NHS Foundation Trust, Birmingham, UK; <sup>3</sup>Centre for Liver & Gastrointestinal Research, Institute of Immunology and Immunotherapy, University of Birmingham, Birmingham, UK; <sup>4</sup>Division of Infectious Diseases and Immunology, Fondazione IRCCS Policlinico San Matteo, University of Pavia, Pavia, Italy; <sup>5</sup>Clinic for Infectious Diseases and Febrile Illnesses, University Medical Centre Ljubljana, Slovenia; <sup>6</sup>Faculty of Medicine, University of Ljubljana, Slovenia; <sup>7</sup>Department of Medicine, University of Seville. Clinical Unit of Infectious Diseases University Hospital Virgen del Rocío. Institute of Biomedicine, Sevilla, CSIC, Spain; <sup>8</sup>Department of Gastroenterology, Hepatology and Endocrinology, Hannover Medical School, Hannover, Germany; <sup>9</sup>Centre for Individualised Infection Medicine (CIIM), Hannover, Germany; <sup>10</sup>Division of Hepatology, Clinic and Polyclinic for Gastroenterology, Hepatology, Infectious Diseases, and Pneumology, University Hospital Leipzig, Leipzig, Germany.*

\*Corresponding author. Address: University Hospital Leipzig, Department of Internal Medicine, Liebigstraße 18, D-04103 Leipzig, Germany; Tel.: +49 341 / 97-12200, fax: +49 341 / 97-12209. E-mail address: [Thomas.Berg@medizin.uni-leipzig.de](mailto:Thomas.Berg@medizin.uni-leipzig.de) (T. Berg).

## Summary

The coronavirus disease 2019 (COVID-19) pandemic poses an enormous challenge to healthcare systems in affected communities. Older patients and those with pre-existing medical conditions have been identified as populations at risk of a severe disease course. It remains unclear at this point to what extent chronic liver diseases should be considered as risk factors, due to a shortage of appropriate studies. However, patients with advanced liver disease and those after liver transplantation represent vulnerable patient cohorts with an increased risk of infection and/or a severe course of COVID-19. In addition, the current pandemic requires unusual allocation of healthcare resources which may negatively impact the care of patients with chronic liver disease that continue to require medical attention. Thus, the challenge hepatologists are facing is to promote telemedicine in the outpatient setting, prioritise outpatient contacts, avoid nosocomial dissemination of the virus to patients and healthcare providers, and at the same time maintain standard care for patients who require immediate medical attention.

## **Introduction**

Coronavirus disease 2019 (COVID-19) is caused by the recently identified coronavirus, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). Symptoms associated with COVID-19 are mostly fever, tiredness and dry cough. Some patients also develop symptoms including shortness of breath, aches, nasal congestion, sore throat, anosmia and ageusia. Nausea, vomiting and diarrhoea occur less frequently [1]. While symptoms remain mild throughout the course of infection in the majority of patients, older patients were identified to be at higher risk of a fatal disease course, with hypertension, diabetes and coronary heart disease the most frequent comorbidities in these cohorts [2, 3].

## **COVID-19 and the liver**

While pre-existing liver disease is not specifically listed in the published cohort studies, elevated alanine aminotransferase (ALT) levels, reduced platelet counts and reduced albumin levels at the time of admission have been associated with higher mortality [2], although not all of these alterations are independent risk factors. It remains unclear at this time whether these laboratory test alterations are a sign for pre-existing liver diseases in patients with a more severe course of infection, whether they rather reflect liver damage caused by the virus itself, or whether they mirror a severe inflammatory response (hypoalbuminemia) with disseminated intravascular coagulation (thrombocytopenia). Possibly, patients with advanced chronic liver disease are at increased risk of infection due to cirrhosis-associated immune dysfunction [4]. The same could be true for patients after liver transplantation and possibly those with autoimmune liver diseases who receive immunosuppressive therapies.

However, currently, there are only limited data available linking underlying liver diseases with the course of SARS-CoV-2 infection and there are many open questions (Box 1). Chronic hepatitis B virus infection for example, which is more prevalent in China than Europe, does not appear to affect the outcome of COVID-19 [1]. In addition, although there is no proof for this assumption, immunosuppression may even provide some protection from immunopathology, which appears to contribute to lung damage in cases with more severe

manifestations of the disease [5, 6]. These may be associated with a macrophage activation syndrome in the context of a hyperinflammatory syndrome characterised by a cytokine storm with multi-organ failure [7]. On the other hand, systemic viral infections are often associated with transient elevations of transaminases which may reflect general immune activation or inflammation caused by circulating cytokines without compromising liver function, a phenomenon called “bystander hepatitis”. This may also be the case in patients with COVID-19 where liver failure has not been specifically reported, even in the most severe and fatal courses of disease [1, 2]. However, signs of liver dysfunction may occur in critically ill patients with COVID-19 [8], as reviewed in [9, 10]. Whether patients with cirrhosis and COVID-19 are at increased risk of decompensation or development of acute-on-chronic liver failure (ACLF), as has been shown for influenza infection [11], remains to be determined. SARS-CoV-2 may also directly infect liver cells as the receptor of the virus, angiotensin-converting enzyme 2 (ACE2) [12], has been shown to be expressed on cholangiocytes [13, 14], which may explain cases of viral shedding into the faeces [15, 16]. How far observations from Asian cohorts can be transferred to European populations remains to be determined, but it appears unlikely that SARS-CoV-2 infection causes liver damage to an amount that substantially contributes to the overall disease burden.

Aside from the direct implications of liver disease on the course of COVID-19 or vice versa, there are several additional aspects that require attention. The current pandemic and the amount of available information – including misinformation and rumours – result in uncertainties and concerns among patients but also healthcare providers. While the threat that COVID-19 poses should not be underestimated, it remains important to maintain care of patients with chronic liver disease and to identify potential ways to prioritise care of these patients in times of limited healthcare resources.

**Box 1. Open questions for liver related basic/translational research regarding COVID-19.**

- Hepatobiliary and gastrointestinal tracts as a hub for viral trafficking via liver and biliary excretion into the gastrointestinal tract
- Influence of immunosuppression on course of COVID-19
- Cirrhosis-associated immune dysfunction and emergence of protective immunity
- Innate and adaptive immune activation or exhaustion in COVID-19 and their impact on chronic liver disease
- Risk of SARS-CoV-2 induced hepatic decompensation/ACLF or graft rejection

ACLF, acute-on-chronic liver failure; COVID-19, coronavirus disease 2019; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2.

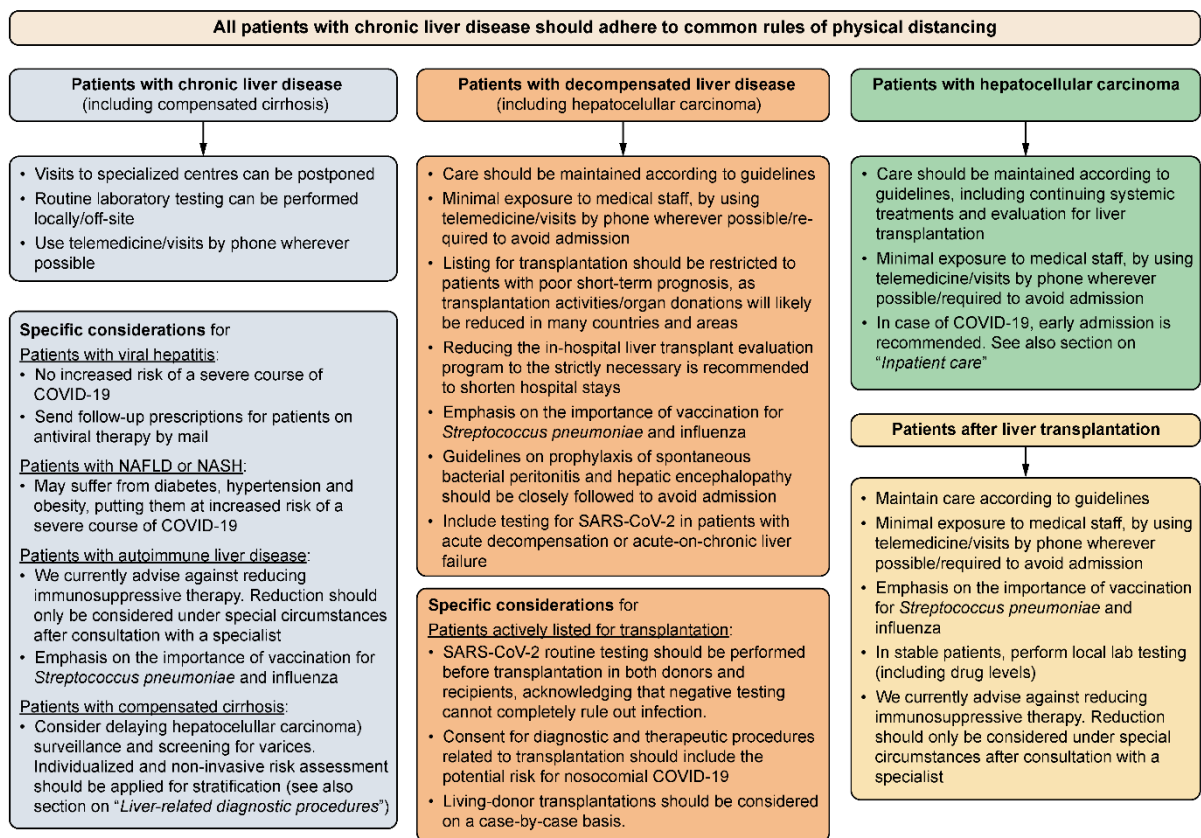
**Considerations for patients with chronic liver disease and patients after liver transplantation**

The recommendations provided here address the specific characteristics of patients with liver disease and are meant to provide additional guidance for the care of these patients. General recommendations and guidelines with regards to prevention, diagnosis and treatment of COVID-19 from local authorities should be adhered to.

**Outpatient care**

The management and surveillance of patients with advanced liver disease and those receiving immunosuppressive treatment is often performed in larger units or centres. These institutions, however, are currently also COVID-19 hotspots, thus, potentially putting outpatients with chronic liver diseases at risk of nosocomial infections. In addition, hospital staff face challenges such as long working hours and even reduced staffing because of COVID-19 quarantining. Therefore, several factors have to be considered by hepatologists who provide care for these vulnerable patients. Clearly, prioritisation criteria need to be defined for outpatient contacts. The general and specific recommendations provided here (Fig. 1) cannot comprehensively cover all patient cohorts and are not backed up by datasets. Moreover, the precise management of these patients strongly depends on the local COVID-19 burden. In case of COVID-19 in patients with more advanced chronic liver disease, we recommend admission for

inpatient care depending on the presence of certain risk factors, which is discussed in more detail in a separate section.



**Fig. 1. Flow chart for the prioritisation of patient care in patients with chronic liver disease.** The individual management of these patients strongly depends on the local COVID-19 burden and officially implemented rules and regulations. In some countries and areas, maintenance of standard care might not be able and transplantation activities might be reduced. COVID-19, coronavirus disease 2019; HCC, hepatocellular carcinoma; NAFLD, non-alcoholic fatty liver disease; NASH, non-alcoholic steatohepatitis; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2.

## Reduction of direct exposure

Physical distancing has been suggested and/or officially implemented in multiple countries across the globe to prevent dissemination of SARS-CoV-2. This measure aims to prevent a rapid increase in SARS-CoV-2 infections in the general population with the overall goal of minimising the number of patients with a severe course of COVID-19 at any one time, enabling healthcare providers to maintain medical/intensive care of these patients. At the same time, vulnerable patient cohorts should be protected from individuals with potential SARS-CoV-2 exposure or infection. Thus, in these patients, the benefits of maintaining patient care have to be weighed against the risk of infection. These considerations require adaption of standard

operating procedures for outpatient care, including remodelling of waiting areas to allow sufficient distance between patients, reduction of waiting times and encouraging patients to wait outside wherever possible and eventually be contacted by phone. In addition, exposure to medical staff should be minimised wherever possible.

Face-to-face contact is the basis for the physician-patient relationships and allows physicians to quickly assess the overall condition of the patient. However, due to the amount of individual contacts, physicians are at risk of infection and subsequently also dissemination of the virus. Thus, it seems appropriate to also limit the amount of face-to-face contacts for patients at risk of a more severe course of infection. Technical solutions are available to enable remote physician-patient interactions and their benefits have already been discussed in the context of the COVID-19 pandemic [17]. Health authorities should be urged to equip hospitals with such systems not only to attend to COVID-19 patients who stay in quarantine at home but also to care for patients that need to be protected from a potentially harmful infection and guide them through the pandemic, explaining the future treatment/management strategies/plan and provide advice concerning general prevention measures. The current pandemic clearly underlines the importance of making these technical solutions available to health care providers in order to be better prepared for situations like these in the future.

### **Patients with compensated liver disease**

#### General considerations:

- Visit at specialised centres can be postponed.
- Use telemedicine visits by phone wherever possible.
- Routine laboratory testing can be performed locally, e.g. through primary care physician, and its frequency needs careful individual risk-benefit considerations.

#### Specific considerations:

- Chronic viral hepatitis does not appear to increase the risk of a severe course of COVID-19 [1]. Use telemedicine/local laboratory testing for follow-up visits in patients under antiviral therapy, send follow-up-prescriptions by mail.
- Patients with non-alcoholic fatty liver disease (NAFLD) or steatohepatitis (NASH) may suffer from metabolic comorbidities such as diabetes, hypertension and obesity putting them at increased risk of a severe course of COVID-19.
- In patients with autoimmune liver disease, we currently advise against reducing immunosuppressive therapy. Reductions should only be considered under special circumstances (e.g. medication-induced lymphopenia, or bacterial/fungal superinfection in case of severe COVID-19) after consultation of a specialist.
- Emphasis on the importance of vaccination for *Streptococcus pneumoniae* and influenza.
- In patients with compensated cirrhosis, consider delaying hepatocellular carcinoma (HCC) surveillance and screening for varices. Non-invasive risk assessment for the presence of varices should be applied for stratification (thrombocyte count or Baveno VI) [18]. See also section on “liver-related diagnostic procedures”.

## **Patients with decompensated liver disease**

### General considerations:

- Care should be maintained according to guidelines but consider minimal exposure to medical staff, by using telemedicine/visits by phone wherever possible/required to avoid admission.
- Listing for transplantation should be restricted to patients with poor short-term prognosis including those with acute/acute-on-chronic liver failure (ALF/ACLF), high model for end-stage liver disease (MELD) score (including exceptional MELDs) and HCC at the upper limits of the Milan criteria, as transplantation activities/organ donations will likely be reduced in many countries and areas.
- Reducing the in-hospital liver transplant evaluation program to the strictly necessary is recommended to shorten the in-hospital stay and also reduce the number of consultations



in other departments/clinics (*i.e.* ophthalmologic, dermatologic, dental, neurologic consultations can be done in local outpatient settings).

- Emphasis on the importance of vaccination for *Streptococcus pneumoniae* and influenza.
- Guidelines on prophylaxis of spontaneous bacterial peritonitis and hepatic encephalopathy should be closely followed to prevent decompensation and avoid admission.
- Include testing for SARS-CoV-2 in patients with acute decompensation or ACLF.

#### Specific considerations for patients actively listed for transplantation:

- SARS-CoV-2 routine testing should be performed before transplantation in both donors and recipients, acknowledging that negative testing cannot completely rule out infection.
- Consent for diagnostic and therapeutic procedures related to transplantation should include the potential risk of nosocomial COVID-19.
- Living-donor transplantations should be considered on a case-by-case basis.

#### **Patients with hepatocellular carcinoma**

- Care should be maintained according to guidelines including continuing systemic treatments and evaluation for liver transplantation, but consider minimal exposure to medical staff, by using telemedicine/visits by phone wherever possible/required to avoid admission.
- In case of COVID-19, early admission is recommended. See also section on “inpatient care”.

#### **Patients after liver transplantation**

##### General considerations:

- Maintain care according to guidelines, but consider minimal exposure to medical staff, by using telemedicine/visits by phone wherever possible/required to avoid admission.
- Emphasis on the importance of vaccination for *Streptococcus pneumoniae* and influenza.
- In stable patients: perform local laboratory testing (including drug levels).

- We currently advise against reduction of immunosuppressive therapy. Reduction should only be considered under special circumstances (e.g. medication-induced lymphopenia, or bacterial/fungal superinfection in case of severe COVID-19) after consultation of a specialist [19].

## **Liver-related diagnostic procedures**

### Endoscopy

*Patients without COVID-19:* Depending on available resources, screening for varices by esophagogastroduodenoscopy (EGD) should be reserved for patients at risk of variceal bleeding, such as patients with a history of variceal bleeding or signs of significant portal hypertension (ascites, thrombocyte count <100,000/μl). Non-invasive risk assessment for the presence of varices should be applied for stratification (thrombocyte count or Baveno VI).

Endoscopic retrograde cholangiography (ERC) for dilatation or stent replacement in patients after liver transplantation or patients with primary sclerosing cholangitis should be performed after careful individual risk-benefit considerations, including risk for nosocomial SARS-CoV-2 infection depending on local COVID-19 burden.

*Patients with COVID-19:* Endoscopic procedures are associated with an increased risk of disseminating SARS-CoV-2. During EGD or ERC, spreading of virus-containing droplets can occur. In addition, shedding of the virus in the faeces increases the risk of dissemination during colonoscopy. Thus, indications for endoscopic procedures in patients with COVID-19 should be limited to emergencies such as gastrointestinal bleeding, bacterial cholangitis or other life-threatening conditions.

### Ultrasound (HCC surveillance)

*Patients without COVID-19:* HCC surveillance can be deferred based on available resources (including availability of therapeutic options in case of HCC diagnosis) at the centre and the individual risk assessment. Patients with increased risk, such as patients with elevated alpha-

fetoprotein levels, advanced cirrhosis, chronic hepatitis B, NASH/diabetes *etc.*, may be prioritised if resources are limited.

*Patients with COVID-19:* HCC surveillance should be deferred until after recovery.

### Liver biopsy

*Patients without COVID-19:* Recommendations strongly depend on the local COVID-19 burden and the individual indication for histological assessment.

Biopsy should be deferred in patients with

- NAFLD or chronic viral hepatitis (for grading/staging)
- mildly elevated transaminases (*e.g.* ALT <3x the upper limit of normal [ULN]) of unknown aetiology

Biopsy should be performed in patients with

- strongly elevated transaminases (*e.g.* > ALT >5x ULN) of unknown aetiology (in case of suspected autoimmune liver disease, treatment without histological diagnosis can be considered based on individual risk-benefit considerations)
- liver masses suspicious of malignancy

*Patients with COVID-19:* Liver biopsy should be deferred in most patients as

- treatment/care for COVID-19 outweighs diagnosis of co-existing liver disease
- systemic inflammation associated with COVID-19 will obscure aetiology-specific histologic characteristics
- liver biopsy may represent a risk for viral transmission (although the virus has so far not been detected in liver tissue, the expression of its receptor on cholangiocytes suggests that the virus might be present in the liver as well).

### **Collaboration with local health care providers and primary care physicians**

The necessity to reduce travel and limit visits in specialised centres is a challenge for liver disease patients and local health care providers alike. In order to facilitate decentralised care for these patients, close collaborations between specialised hepatologists and local health care

providers is required. This pertains to several factors such as communication of immunosuppressive target levels in patients after liver transplantation, management of patients with severe ALT elevations of unknown aetiologies and others. Thus, we recommend that specialised centres provide easily accessible contact information for local health care providers to facilitate fast and uncomplicated collaboration for the benefit of patients with liver diseases.

### **Inpatient care**

Many patients with chronic liver disease will continue to require inpatient care during the COVID-19 crisis for decompensation, cholangitis, rejection or other complications. General measures to prevent SARS-CoV-2 exposure and infection will be of utmost importance for these patients. Depending on the local infrastructure, implementation of COVID-19-clean wards or hospitals is warranted. However, based on the local COVID-19-burden, a sharp separation of “clean” and “dirty” wards/hospitals may not be sustainable, although recent evidence provides somewhat reassuring data regarding the absence of environmental contamination with SARS-CoV-2 RNA of inanimate surfaces outside patient rooms [20]. Wherever possible, patients with chronic liver disease requiring inpatient care for non-COVID-19 causes should be admitted to COVID-19-clean wards or hospitals. As these institutions may not be able to provide specialised hepatology-care, similar to our recommendations on outpatient care, we recommend that specialised centres provide easily accessible contact information to facilitate immediate hepatology consultations.

However, due to the contagiousness of the virus, patients with underlying liver disease will become infected and will subsequently require inpatient care for COVID-19. With regards to nosocomial-infections, recent observations from Spain that compared patient characteristics between patients with community-acquired influenza and nosocomial-acquired influenza could not detect a significant difference between these cohorts with regards to underlying chronic liver disease [21]. Whether these observations will also hold up in the context of the COVID-19 pandemic remains to be proven. Until further evidence emerges, we recommend that patients with chronic liver disease and COVID-19 are admitted for inpatient care if they have

additional risk factors for a more severe COVID-19 course like hypertension, diabetes or obesity, cirrhosis, HCC or a post-transplant status.

**Table 1. Selected repurposed drugs that have been suggested/discussed for the treatment of COVID-19 and considerations for patients with liver disease or after liver transplantation.**

Drug	Mechanism of action, rationale for COVID-19	Considerations for patients with liver diseases or after liver transplantation
Remdesivir	<ul style="list-style-type: none"> <li>• NUC/viral RNA polymerase inhibitor (completed phase III for Ebola treatment)</li> <li>• Inhibits SARS-CoV-2 <i>in vitro</i> [24]</li> <li>• Case reports with COVID-19 [26]</li> </ul>	<ul style="list-style-type: none"> <li>• No relevant drug-interactions expected [34]</li> <li>• No experience in liver cirrhosis but a NUC might be safer than other drug classes based on experience with NUCs in chronic hepatitis B and C</li> <li>• Liver toxicity (↑ALT) possible</li> </ul>
Chloroquine/hydroxychloroquine [35] ± azithromycin	<ul style="list-style-type: none"> <li>• Interference with the cellular receptor ACE2 [24]</li> <li>• Endosomal acidification fusion inhibitor</li> <li>• Generally used for treatment of malaria and amoebiasis</li> <li>• <i>In vitro</i> and <i>in vivo</i> data [32, 36]</li> </ul>	<ul style="list-style-type: none"> <li>• Exclude G6PD deficiency before application</li> <li>• Drug-interactions with immunosuppressive drugs: close monitoring of drug level is required for cyclosporine, tacrolimus, sirolimus, everolimus [34]</li> <li>• Hydroxychloroquine therapy has not been associated with ALT abnormalities and is an extremely rare cause of clinically apparent acute liver injury (LiverTox).</li> </ul>
Lopinavir/ritonavir	<ul style="list-style-type: none"> <li>• Lopinavir/ritonavir are approved PIs for HIV</li> <li>• <i>In vitro</i> data, experience in patients with SARS, case reports with COVID-19 [37]</li> <li>• No proven efficacy <i>in vivo</i> in severe COVID-19 [28]</li> <li>• Many centres have discontinued its use</li> </ul>	<ul style="list-style-type: none"> <li>• Known and well-studied drug-interactions with immunosuppressive drugs. mTOR inhibitors (sirolimus, everolimus) should not be co-administered, close monitoring of drug level are required for calcineurin-inhibitors (cyclosporin, tacrolimus) [34]</li> <li>• Data for patients with liver cirrhosis exist [38]</li> <li>• The risk of lopinavir-associated hepatotoxicity in patients with very advanced liver disease is low</li> <li>• Based on experience with PIs in HCV, patients with decompensated cirrhosis should not be treated</li> </ul>
Tocilizumab	<ul style="list-style-type: none"> <li>• Humanised mAb targeting interleukin-6 receptor</li> <li>• Treat cytokine release syndrome observed in COVID-19 [7]</li> </ul>	<ul style="list-style-type: none"> <li>• ALT elevations are frequent but clinically apparent liver injury with jaundice seem to be rare [39]</li> <li>• Patients with decompensated cirrhosis should not be treated</li> <li>• Consider risk of HBV reactivation [40]</li> </ul>
Methylprednisolone (steroids)	<ul style="list-style-type: none"> <li>• Corticosteroids bind nuclear receptors to dampen proinflammatory cytokines</li> <li>• Mostly used in patients with septic shock</li> <li>• Currently NOT recommended by WHO [41]</li> </ul>	<ul style="list-style-type: none"> <li>• The risk of other infections (e.g. SBP) and viral shedding may increase in patients with decompensated liver cirrhosis</li> <li>• Consider antimicrobial prophylaxis</li> <li>• Consider risk of HBV reactivation</li> </ul>
Convalescent plasma	<ul style="list-style-type: none"> <li>• Case reports with COVID-19 [42]</li> </ul>	<ul style="list-style-type: none"> <li>• No experience in patients with chronic liver disease</li> </ul>
Umifenovir (Arbidol)*	<ul style="list-style-type: none"> <li>• May inhibit viral entry into target cells and stimulate the immune response, used to treat influenza in some countries [31]</li> </ul>	<ul style="list-style-type: none"> <li>• Possible drug interactions between arbidol and CYP3A4 inhibitors and inducers [43]</li> <li>• Potentially metabolised in liver and intestines in humans. Caution in patients with liver cirrhosis</li> </ul>
Favipiravir/favilavir*	<ul style="list-style-type: none"> <li>• Guanine analogue, RNA-dependent RNA polymerase (RdRp)-inhibitor, approved for influenza in Japan</li> <li>• Preliminary results from a study with 80 COVID-19 patients [31]</li> </ul>	<ul style="list-style-type: none"> <li>• Metabolised by aldehyde oxidase and xanthine oxidase. CYP450 isoenzymes are not involved in the metabolism</li> <li>• Elevation of ALT and AST possible.</li> <li>• No data in cirrhosis available</li> </ul>
Sofosbuvir* <i>maybe combination with ribavirin</i>	<ul style="list-style-type: none"> <li>• Nucleotide analogue, RdRp-inhibitor</li> <li>• Approved for treatment of chronic hepatitis C</li> <li>• <i>In vitro</i> data show binding to SARS-CoV-2 RdRp [44]</li> </ul>	<ul style="list-style-type: none"> <li>• Good experience in patients with chronic hepatitis C including patients with decompensated cirrhosis</li> <li>• For drug-interaction details see [45]</li> <li>• Ribavirin may cause severe haemolytic anaemia</li> </ul>

Baricitinib*	<ul style="list-style-type: none"> <li>• Janus kinase inhibitor, might interrupt endocytosis of the virus and intracellular assembly of virus particles [46]</li> <li>• Could affect both inflammation and cellular viral entry</li> </ul>	<ul style="list-style-type: none"> <li>• Associated with transient and usually mild elevations of ALT [47]</li> <li>• Patients with decompensated cirrhosis should not be treated</li> </ul>
Camostat*	<ul style="list-style-type: none"> <li>• Blocks serine protease TMPRSS2 <i>in vitro</i> which is required for S protein priming [48]</li> <li>• Licensed in Japan for treatment of chronic pancreatitis</li> </ul>	<ul style="list-style-type: none"> <li>• Patients with chronic viral hepatitis and cirrhosis are excluded from clinical trial for chronic pancreatitis [49].</li> <li>• Drug-interactions unknown</li> </ul>
Emapalumab*	<ul style="list-style-type: none"> <li>• mAb targeting interferon-gamma</li> <li>• Treat cytokine release syndrome observed in COVID-19 [7]</li> <li>• Approved for haemophagocytic lymphohistiocytosis</li> <li>• Clinical trials for COVID-19 planned [50]</li> </ul>	<ul style="list-style-type: none"> <li>• Associated with mild and transient ALT elevations typically arising a few weeks after start of treatment</li> <li>• Risk of reactivation of tuberculosis, pneumocystis jirovecii, herpes zoster</li> <li>• Risk of HBV reactivation may be lower</li> </ul>
Anakinra*	<ul style="list-style-type: none"> <li>• Interleukin 1 receptor antagonist</li> <li>• Clinical trials for COVID-19 planned [50]</li> </ul>	<ul style="list-style-type: none"> <li>• Minimal hepatic metabolism</li> </ul>

ACE2, angiotensin-converting enzyme 2; ALT, alanine aminotransferase; AST, aspartate aminotransferase; COVID-19, coronavirus disease 2019; NUC, nucleoside analogue; PIs, protease inhibitors; RdRp, RNA-dependent RNA polymerase; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2.

This list is not intended to give treatment recommendations. The evidence to use these drugs is low. Multiple RCTs for the above-mentioned drugs and some other drugs are ongoing in patients with SARS-CoV-2 infection.

\*These drugs are listed to include the most frequently discussed compounds for COVID-19. These drugs may lack clinical efficacy, and some have been considered based on *in vitro* observations or their mode of action.

## Treatment considerations for COVID-19

Although there are currently no drugs approved for SARS-CoV-2, several repurposed drugs have been tested in recent weeks and many of them are still under investigation (see Table 1) [22]. Remdesivir acts as an adenosine-analogue that induces RNA chain termination and was initially developed as an antiviral agent against Ebola [23]. It has recently been shown to inhibit a clinical isolate of SARS-CoV-2 *in vitro* [24] and to reduce disease severity of the related MERS-CoV-infection in a non-human primate model *in vivo* [25]. In addition, several reports have suggested clinical efficacy in patients suffering from COVID-19 [26, 27]. Clinical trials are ongoing to further evaluate efficacy in affected patients. Although ritonavir-boosted lopinavir showed some antiviral effect on SARS-CoV-2 *in vitro*, a recently published clinical trial in patients with severe COVID-19 showed no effect *in vivo* compared to no treatment [28]. Other drugs currently under evaluation include chloroquine phosphate or hydroxychloroquine [29].

Chloroquine has shown antiviral efficacy against SARS-CoV-2 *in vitro* through interference with the ACE2-receptor mediated endocytosis and is widely used in the treatment of patients with severe COVID-19 as monotherapy, but also in combination with azithromycin [24, 30-32]. The administration of any of these compounds to infected patients will remain an individual decision as efficacy and optimal timing remain to be clarified. However, with regards to patients with chronic liver disease, possible adverse events have to be kept in mind. This is particularly important with respect to drug-interactions in patients with certain immunosuppressive therapies where drug levels of cyclosporine, tacrolimus, sirolimus or everolimus will have to be closely monitored. In addition, patients with impaired liver function are at high risk of drug toxicities, *i.e.* in patients with Child-Pugh B/C cirrhosis. Table 1 summarises some considerations with regards to potential toxicities in these patients. It is also important to stress that all the drugs currently under investigation are not approved for SARS-CoV-2. However, given that early initiation of antiviral therapy is known to blunt the course of influenza, it is reasonable to assume that early treatment initiation might also be beneficial to prevent severe pneumonia in COVID-19. Thus, in patients with liver disease and risk factors for a severe course of the disease, we recommend inclusion into early antiviral treatment programmes or clinical trials that might be active at different centres.

#### General considerations for patients with chronic liver disease and COVID-19:

- Consider early admission according to the presence of additional risk factors and inclusion in clinical trials and (experimental) antiviral therapy of COVID-19 following local guidelines (see Table 1 for guidance with regards to CLD)
- Prevent acetaminophen overdosing (2–3 g/day is considered safe in patients with cirrhosis without active alcohol consumption [33])
- Do not administer non-steroidal anti-inflammatory drugs in patients with cirrhosis and portal hypertension [33]
- See also section on “liver-related diagnostic procedures” for recommendations on endoscopy, HCC surveillance and liver biopsy

### Specific considerations for patients with chronic liver disease and COVID-19:

- In patients with decompensated cirrhosis, treatment for cirrhosis-associated complications such as portal hypertension, ascites, hepatic encephalopathy, spontaneous bacterial peritonitis *etc.* should be continued
- In patients with HCC, locoregional therapies should be postponed whenever possible and immune-checkpoint inhibitor therapy temporarily withdrawn. The decision on whether to continue (reduced dose) kinase inhibitors in non-severe COVID-19 should be taken on a case-by-case basis.
- In patients after liver transplantation, dose adjustment of calceurin- and/or mTOR-inhibitors might be required depending on the antiviral therapy initiated (Table 1).

### **Acknowledgement**

We thank Benjamin Maasoumy for critically reviewing the manuscript and providing valuable comments.

### **References**

- [1] Guan W-j, Ni Z-y, Hu Y, Liang W-h, Ou C-q, He J-x, et al. Clinical Characteristics of Coronavirus Disease 2019 in China. *New England Journal of Medicine* 2020.
- [2] Zhou F, Yu T, Du R, Fan G, Liu Y, Liu Z, et al. Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: a retrospective cohort study. *The Lancet* 2020.
- [3] Yang J, Zheng Y, Gou X, Pu K, Chen Z, Guo Q, et al. Prevalence of comorbidities in the novel Wuhan coronavirus (COVID-19) infection: a systematic review and meta-analysis. *International Journal of Infectious Diseases* 2020.
- [4] Albillos A, Lario M, Álvarez-Mon M. Cirrhosis-associated immune dysfunction: Distinctive features and clinical relevance. *Journal of Hepatology* 2014;61:1385-1396.
- [5] Xu Z, Shi L, Wang Y, Zhang J, Huang L, Zhang C, et al. Pathological findings of COVID-19 associated with acute respiratory distress syndrome. *The Lancet Respiratory Medicine* 2020.
- [6] Li X, Geng M, Peng Y, Meng L, Lu S. Molecular immune pathogenesis and diagnosis of COVID-19. *Journal of Pharmaceutical Analysis* 2020.
- [7] Mehta P, McAuley DF, Brown M, Sanchez E, Tattersall RS, Manson JJ. COVID-19: consider cytokine storm syndromes and immunosuppression. *The Lancet* 2020.
- [8] Yang X, Yu Y, Xu J, Shu H, Xia Ja, Liu H, et al. Clinical course and outcomes of critically ill patients with SARS-CoV-2 pneumonia in Wuhan, China: a single-centered, retrospective, observational study. *The Lancet Respiratory Medicine* 2020.
- [9] Zhang C, Shi L, Wang F-S. Liver injury in COVID-19: management and challenges. *The Lancet Gastroenterology & Hepatology* 2020.



- [10] Xu L, Liu J, Lu M, Yang D, Zheng X. Liver injury during highly pathogenic human coronavirus infections. *Liver International* 2020;n/a.
- [11] Schütte A, Ciesek S, Wedemeyer H, Lange CM. Influenza virus infection as precipitating event of acute-on-chronic liver failure. *Journal of Hepatology* 2019;70:797-799.
- [12] Walls AC, Park Y-J, Tortorici MA, Wall A, McGuire AT, Velesler D. Structure, Function, and Antigenicity of the SARS-CoV-2 Spike Glycoprotein. *Cell* 2020.
- [13] Chai X, Hu L, Zhang Y, Han W, Lu Z, Ke A, et al. Specific ACE2 Expression in Cholangiocytes May Cause Liver Damage After 2019-nCoV Infection. *bioRxiv* 2020:2020.2002.2003.931766.
- [14] Xu H, Zhong L, Deng J, Peng J, Dan H, Zeng X, et al. High expression of ACE2 receptor of 2019-nCoV on the epithelial cells of oral mucosa. *International Journal of Oral Science* 2020;12:8.
- [15] Yeo C, Kaushal S, Yeo D. Enteric involvement of coronaviruses: is faecal-oral transmission of SARS-CoV-2 possible? *The Lancet Gastroenterology & Hepatology* 2020;5:335-337.
- [16] Gu J, Han B, Wang J. COVID-19: Gastrointestinal manifestations and potential fecal-oral transmission. *Gastroenterology* 2020.
- [17] Hollander JE, Carr BG. Virtually Perfect? Telemedicine for Covid-19. *New England Journal of Medicine* 2020.
- [18] Maurice JB, Brodtkin E, Arnold F, Navaratnam A, Paine H, Khawar S, et al. Validation of the Baveno VI criteria to identify low risk cirrhotic patients not requiring endoscopic surveillance for varices. *Journal of Hepatology* 2016;65:899-905.
- [19] Zhu L, Xu X, Ma K, Yang J, Guan H, Chen S, et al. Successful recovery of COVID-19 pneumonia in a renal transplant recipient with long-term immunosuppression. *American Journal of Transplantation* 2020;n/a.
- [20] Colaneri M, Seminari E, Piralla A, Zuccaro V, Filippo AD, Baldanti F, et al. Lack of SARS-CoV-2 RNA environmental contamination in a tertiary referral hospital for infectious diseases in Northern Italy. *Journal of Hospital Infection*.
- [21] Godoy P, Torner N, Soldevila N, Rius C, Jane M, Martínez A, et al. Hospital-acquired influenza infections detected by a surveillance system over six seasons, from 2010/2011 to 2015/2016. *BMC Infectious Diseases* 2020;20:80.
- [22] Harrison C. Coronavirus puts drug repurposing on the fast track. 2020 [cited March 12th 2020]; Available from: <https://www.nature.com/articles/d41587-020-00003-1>
- [23] Warren TK, Jordan R, Lo MK, Ray AS, Mackman RL, Soloveva V, et al. Therapeutic efficacy of the small molecule GS-5734 against Ebola virus in rhesus monkeys. *Nature* 2016;531:381-385.
- [24] Wang M, Cao R, Zhang L, Yang X, Liu J, Xu M, et al. Remdesivir and chloroquine effectively inhibit the recently emerged novel coronavirus (2019-nCoV) in vitro. *Cell Research* 2020;30:269-271.
- [25] de Wit E, Feldmann F, Cronin J, Jordan R, Okumura A, Thomas T, et al. Prophylactic and therapeutic remdesivir (GS-5734) treatment in the rhesus macaque model of MERS-CoV infection. *Proceedings of the National Academy of Sciences* 2020:201922083.
- [26] Holshue ML, DeBolt C, Lindquist S, Lofy KH, Wiesman J, Bruce H, et al. First Case of 2019 Novel Coronavirus in the United States. *New England Journal of Medicine* 2020;382:929-936.
- [27] Sheahan TP, Sims AC, Leist SR, Schäfer A, Won J, Brown AJ, et al. Comparative therapeutic efficacy of remdesivir and combination lopinavir, ritonavir, and interferon beta against MERS-CoV. *Nature Communications* 2020;11:222.
- [28] Cao B, Wang Y, Wen D, Liu W, Wang J, Fan G, et al. A Trial of Lopinavir–Ritonavir in Adults Hospitalized with Severe Covid-19. *New England Journal of Medicine* 2020.
- [29] Gao J, Tian Z, Yang X. Breakthrough: Chloroquine phosphate has shown apparent efficacy in treatment of COVID-19 associated pneumonia in clinical studies. *BioScience Trends* 2020;advpub.
- [30] Li G, De Clercq E. Therapeutic options for the 2019 novel coronavirus (2019-nCoV). *Nat Rev Drug Discov* 2020;19:149-150.

- [31] Dong L, Hu S, Gao J. Discovering drugs to treat coronavirus disease 2019 (COVID-19). *Drug Discoveries & Therapeutics* 2020;14:58-60.
- [32] Gautret P, Lagier J-C, Parola P, Hoang VT, Meddeb L, Mailhe M, et al. Hydroxychloroquine and azithromycin as a treatment of COVID-19: results of an open-label non-randomized clinical trial. *International Journal of Antimicrobial Agents* 2020:105949.
- [33] Chandok N, Watt KDS. Pain Management in the Cirrhotic Patient: The Clinical Challenge. *Mayo Clinic Proceedings* 2010;85:451-458.
- [34] COVID-19 Drug interactions. 2020 [cited March 16th 2020]; Available from: <http://www.covid19-druginteractions.org/>
- [35] [Expert consensus on chloroquine phosphate for the treatment of novel coronavirus pneumonia]. *Zhonghua jie he he hu xi za zhi = Zhonghua jiehe he huxi zazhi = Chinese journal of tuberculosis and respiratory diseases* 2020;43:E019.
- [36] Al-Bari MAA. Targeting endosomal acidification by chloroquine analogs as a promising strategy for the treatment of emerging viral diseases. *Pharmacology Research & Perspectives* 2017;5:e00293.
- [37] Young BE, Ong SWX, Kalimuddin S, Low JG, Tan SY, Loh J, et al. Epidemiologic Features and Clinical Course of Patients Infected With SARS-CoV-2 in Singapore. *JAMA* 2020.
- [38] Casado JL, Del Palacio M, Moya J, Rodriguez JM, Moreno A, Perez-Elías MJ, et al. Safety and Pharmacokinetics of Lopinavir in HIV/HCV Coinfected Patients with Advanced Liver Disease. *HIV Clinical Trials* 2011;12:235-243.
- [39] Genovese MC, Kremer JM, van Vollenhoven RF, Alten R, Scali JJ, Kelman A, et al. Transaminase Levels and Hepatic Events During Tocilizumab Treatment: Pooled Analysis of Long-Term Clinical Trial Safety Data in Rheumatoid Arthritis. *Arthritis & Rheumatology* 2017;69:1751-1761.
- [40] Chen L-F, Mo Y-Q, Jing J, Ma J-D, Zheng D-H, Dai L. Short-course tocilizumab increases risk of hepatitis B virus reactivation in patients with rheumatoid arthritis: a prospective clinical observation. *International Journal of Rheumatic Diseases* 2017;20:859-869.
- [41] WHO. Clinical management of severe acute respiratory infection (SARI) when COVID-19 disease is suspected. 2020 [cited; Available from: [https://www.who.int/publications-detail/clinical-management-of-severe-acute-respiratory-infection-when-novel-coronavirus-\(ncov\)-infection-is-suspected](https://www.who.int/publications-detail/clinical-management-of-severe-acute-respiratory-infection-when-novel-coronavirus-(ncov)-infection-is-suspected)
- [42] Shen C, Wang Z, Zhao F, Yang Y, Li J, Yuan J, et al. Treatment of 5 Critically Ill Patients With COVID-19 With Convalescent Plasma. *JAMA* 2020.
- [43] Deng P, Zhong D, Yu K, Zhang Y, Wang T, Chen X. Pharmacokinetics, Metabolism, and Excretion of the Antiviral Drug Arbidol in Humans. *Antimicrobial Agents and Chemotherapy* 2013;57:1743-1755.
- [44] Elfiky AA. Anti-HCV, nucleotide inhibitors, repurposing against COVID-19. *Life Sciences* 2020;248:117477.
- [45] HEP Drug interactions. 2020 [cited March 18th 2020]; Available from: <https://www.hep-druginteractions.org/>
- [46] Richardson P, Griffin I, Tucker C, Smith D, Oechsle O, Phelan A, et al. Baricitinib as potential treatment for 2019-nCoV acute respiratory disease. *The Lancet* 2020;395:e30-e31.
- [47] Markham A. Baricitinib: First Global Approval. *Drugs* 2017;77:697-704.
- [48] Hoffmann M, Kleine-Weber H, Schroeder S, Krüger N, Herrler T, Erichsen S, et al. SARS-CoV-2 Cell Entry Depends on ACE2 and TMPRSS2 and Is Blocked by a Clinically Proven Protease Inhibitor. *Cell* 2020.
- [49] Ramsey ML, Nuttall J, Hart PA, on behalf of the TIT. A phase 1/2 trial to evaluate the pharmacokinetics, safety, and efficacy of NI-03 in patients with chronic pancreatitis: study protocol for a randomized controlled trial on the assessment of camostat treatment in chronic pancreatitis (TACTIC). *Trials* 2019;20:501.

- [50] Press release on planned clinical trials to evaluate anakinra and emapalumab in COVID-19. 2020 [cited March 30th, 2020]; Available from: <https://www.sobi.com/en/press-releases/sobi-initiate-clinical-study-evaluate-whether-anakinra-and-emapalumab-may-relieve>
- [51] van Doremalen N, Bushmaker T, Morris DH, Holbrook MG, Gamble A, Williamson BN, et al. Aerosol and Surface Stability of SARS-CoV-2 as Compared with SARS-CoV-1. *New England Journal of Medicine* 2020.
- [52] Ghinai I, McPherson TD, Hunter JC, Kirking HL, Christiansen D, Joshi K, et al. First known person-to-person transmission of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) in the USA. *The Lancet* 2020.
- [53] Liu Y, Gayle AA, Wilder-Smith A, Rocklöv J. The reproductive number of COVID-19 is higher compared to SARS coronavirus. *Journal of Travel Medicine* 2020;27.
- [54] Wang W, Tang J, Wei F. Updated understanding of the outbreak of 2019 novel coronavirus (2019-nCoV) in Wuhan, China. *Journal of Medical Virology* 2020;92:441-447.
- [55] Backer JA, Klinkenberg D, Wallinga J. Incubation period of 2019 novel coronavirus (2019-nCoV) infections among travellers from Wuhan, China, 20–28 January 2020. *Eurosurveillance* 2020;25:2000062.
- [56] Fang Y, Zhang H, Xie J, Lin M, Ying L, Pang P, et al. Sensitivity of Chest CT for COVID-19: Comparison to RT-PCR. *Radiology*;0:200432.
- [57] Adhikari SP, Meng S, Wu Y-J, Mao Y-P, Ye R-X, Wang Q-Z, et al. Epidemiology, causes, clinical manifestation and diagnosis, prevention and control of coronavirus disease (COVID-19) during the early outbreak period: a scoping review. *Infectious Diseases of Poverty* 2020;9:29.
- [58] Hellewell J, Abbott S, Gimma A, Bosse NI, Jarvis CI, Russell TW, et al. Feasibility of controlling COVID-19 outbreaks by isolation of cases and contacts. *The Lancet Global Health* 2020;8:e488-e496.
- [59] Park GE, Ko J-H, Peck KR, Lee JY, Lee JY, Cho SY, et al. Control of an Outbreak of Middle East Respiratory Syndrome in a Tertiary Hospital in Korea. *Annals of Internal Medicine* 2016;165:87-93.
- [60] Klompas M. Coronavirus Disease 2019 (COVID-19): Protecting Hospitals From the Invisible. *Annals of Internal Medicine* 2020.
- [61] Thomas RF, Christopher TL. Identifying and Interrupting Superspreading Events—Implications for Control of Severe Acute Respiratory Syndrome Coronavirus 2. *Emerging Infectious Disease journal* 2020;26.
- [62] Ong SWX, Tan YK, Chia PY, Lee TH, Ng OT, Wong MSY, et al. Air, Surface Environmental, and Personal Protective Equipment Contamination by Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) From a Symptomatic Patient. *JAMA* 2020.
- [63] Kuster GM, Pfister O, Burkard T, Zhou Q, Twerenbold R, Haaf P, et al. SARS-CoV2: should inhibitors of the renin–angiotensin system be withdrawn in patients with COVID-19? *European Heart Journal* 2020.
- [64] Position Statement of the ESC Council on Hypertension on ACE-Inhibitors and Angiotensin Receptor Blockers. 2020 [cited March 30th, 2020]; Available from: [https://www.escardio.org/Councils/Council-on-Hypertension-\(CHT\)/News/position-statement-of-the-esc-council-on-hypertension-on-ace-inhibitors-and-ang](https://www.escardio.org/Councils/Council-on-Hypertension-(CHT)/News/position-statement-of-the-esc-council-on-hypertension-on-ace-inhibitors-and-ang)

## **Appendix on general recommendations for the prevention of SARS-CoV-2 infection**

Even though our understanding of the epidemiology of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection remains limited, 3 methods of virus transmission have been reported: person-to-person transmission through respiratory droplets, contact transmission by touching contaminated surfaces followed by touching mouths, nose or eyes, and aerosol transmission by inhaling high concentrations of foaming aerosols in a relatively closed environment, which happens on specific occasions [51, 52]. In addition, the digestive system has been indicated as a possible route of viral transmission, as discussed above. The transmission of SARS-CoV-2 is often unpredictable and difficult to prevent. Therefore, understanding the viral, host, environmental, and behavioural characteristics may help in creating strategies for prevention and control of viral transmission.

Studies have shown that SARS-CoV-2 has a high level of transmissibility. The estimated initial median of the basic reproduction ( $R_0$ ) was shown to be 2.79, meaning that one infected person will – on average – infect 2.79 other people, however  $R_0$  differs among various regions [53]. Besides, even asymptotically infected persons or those presenting with minimal symptoms were found to be contagious [54]. Reports have already shown that the incubation period can be extended beyond 14 days, however, the vast majority of cases becomes symptomatic within 14 days [55]. The signs and symptoms of coronavirus disease 2019 (COVID-19) are not specific and are therefore difficult to distinguish from those of other respiratory virus infections. The 70% sensitivity of a single nasopharyngeal swab performed early in the course of disease represents an additional obstacle in making an accurate diagnosis [56]. Delayed diagnoses were reported to be one of the leading reasons for nosocomial transmissions.

Thus, effective prevention and control measures that are rapidly implemented and highly targeted within the community and healthcare settings are of critical importance to prevent transmission of SARS-CoV-2. For the general population, behavioural recommendations such as washing hands, covering coughs, and social distance are needed, whereas for healthcare workers extremely rigorous infection control has to be implemented

[57]. Prompt identification of infected persons, their isolation and contact tracing might be sufficient to reduce viral transmission allowing control of a cluster of infected individuals. However, delays in isolation of patients from the onset of symptoms will markedly reduce such control [58].

Healthcare settings represent one of the most critical environments for prevention and control of SARS-CoV-2 transmission. Universal infection control procedures are necessary to be performed in all areas of all settings. Besides, precise triage protocols, special behavioural protocols and patterns for staff, patients, and visitors, rapid diagnosis and strict isolation such as isolation rooms, partitions to protect against respiratory droplets and ventilation systems must be implemented immediately [59, 60]. Measures are targeted to enable rapid detection and isolation of all potentially contagious patients and those with high suspicion of being infected, and to prevent the further spread of infection.

Special attention must be paid to infection control during certain procedures which enable aerosolization of virus-containing droplets, producing a much more effectively and widely disseminated infection. Such procedures include bronchoscopy, intubation, suctioning, sputum induction, nebulizer therapies, as well as esophagogastroduodenoscopy [61]. They should be performed under strict infection control procedures and, if possible, take place in units with airborne infection isolation. Although the virus has so far not been detected in liver tissue, the expression of its receptor on cholangiocytes and shedding of the virus in the faeces [16] suggest that the virus might be present in the liver as well. Thus, liver biopsy and certainly colonoscopy may also represent a risk for viral transmission if strict infection control measures are not implemented. Basically, ensuring cleanliness of toilets and other potentially contaminated surfaces is of extreme importance. Effective decontamination and strict environmental hygiene are crucial in elimination of respiratory droplets and faecal shedding from a contaminated healthcare environment [62]. Finally, in case of nosocomial transmission, possible modes of spread have to be carefully analysed, to improve strategies for further prevention of viral transmission.

It has recently been suggested that several drugs could increase susceptibility for SARS-CoV-2 infection and/or result in a more severe course of COVID-19. Most prominently, Ibuprofen and angiotensin-converting enzyme (ACE)-inhibitors have been discussed. While for Ibuprofen no actual or theoretical explanation has been proposed, ACE-I have been suggested to upregulate ACE2, which could increase susceptibility for SARS-CoV-2. Based on this hypothesis, discontinuation of ACE-I usage in patients with hypertension and chronic heart disease has been discussed. However, based on the sparse evidence that is currently available, ACE-I should not be withdrawn in patients on stable medication until further evidence emerges [63, 64].