



COVID-19: Thoughts and comments from a tertiary liver transplant center in France

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► **To cite this version:**

Stylios Tzedakis, Heithem Jeddou, Pauline Houssel-Debry, Laurent Sulpice, Karim Boudjema. COVID-19: Thoughts and comments from a tertiary liver transplant center in France. *American Journal of Transplantation*, Wiley, 2020, 20 (7), pp.1952-1953. 10.1111/ajt.15918 . hal-02798003

HAL Id: hal-02798003

<https://hal-univ-rennes1.archives-ouvertes.fr/hal-02798003>

Submitted on 15 Jun 2020

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Article type : Letter to the Editor

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COVID-19: Thoughts and comments from a Tertiary Liver Transplant Center in France

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ABBREVIATIONS

ACE2, Angiotensin converting enzyme 2

BAL, Bronchoalveolar lavage

COVID-19, Coronavirus disease 2019

ECDC, European Centre for Disease Prevention and Control

ICU, Intensive Care Unit

MELD, Model for end-stage liver disease

MERS-CoV, Middle East respiratory syndrome - coronavirus

NPS, Nasopharyngeal swab

SARS-CoV, Severe acute respiratory syndrome – coronavirus

We read with interest the analysis on modern approaches to organ transplantation during the COVID-19 pandemic by Kumar et al.¹. Following the December 2019 SARS-CoV-2 outbreak in China, France is now fifth in number of deaths worldwide while at the same time hosting one of the biggest European liver transplantation programs.

Past experience with other Coronaviruses outbreaks such as the SARS-CoV and MERS-CoV has shown possible transplant recipient contamination and death through their ability to bind to the ACE2 receptors essentially located in the upper and lower respiratory tract but also in liver and bile duct cells²⁻⁴. Moreover, liver transplant recipients seem to have more prolonged shedding of virus, owing to immunosuppression, thus potentially increasing the risk of transmission to contacts (super-spreader's effect)².

Our hospital, located in Rennes, west France (Brittany), hosts the second biggest national liver transplant center. As liver transplant specialists, we think it is crucial to carefully balance costs and benefits in performing a liver transplantation within a COVID-19 epidemic region during a COVID-19 outbreak and decreasing transplant activity might be needed (e.g. local center activity has decreased by 60% since the pandemic outbreak).

We thought useful to share French experience and propositions for liver transplantation. In accordance with ECDC and Kumar et al., donors who are infected with COVID-19 are not eligible for organ donation, as risk for donor-recipient transmission cannot be reliably evaluated. All liver transplant donors and candidates (as opposed to selective screening referred by Kumar et al.) without symptoms or diagnosis of COVID-19 in an epidemic area should be tested for the presence of SARS-CoV-2 in the BAL or NPS specimens collected before validating organ procurement and transplant procedure (results taking up to 6 hours) in order to assure recipient's and organ procurement teams' safety. In France, all deceased donors systematically undergo a full-body scan (adding to the discussion made by Kumar et al.) which may help early COVID diagnosis⁵. Liver transplantation is currently reserved to the most urgent cases, which means candidates with fulminant hepatic failure, high MELD score (> 25) or for which the 3-month risk of 'Drop-out' (e.g. malignant disease progression) is greater than the risk of getting infected with SARS-CoV-2. These 'urgent' candidates are in most cases in-hospital patients, COVID testing is performed early and transplant timing is not impacted. Local ICU capacities play a pivotal role and full bed occupation may impose a total cease of activity. Liver transplant candidates diagnosed with COVID-19 (an issue not addressed by Kumar et al.) are put in temporary contraindication for liver transplantation and are re-tested 8 days later. If NPS or BAL becomes negative, liver transplantation is permitted. Early transplant recipients diagnosed with COVID-19 during the postoperative course are transferred to SARS-CoV-2 specialized units in order to protect hospital personnel. Finally, all liver transplant recipients should be informed of the current

epidemic outbreak, possibilities for transplantation and should be excluded from all COVID-independent clinical research studies they may have consent to participate as inclusion of infected patients may bias study outcomes.

Disclosures: The authors of this manuscript have no conflicts of interest to disclose as described by the *American Journal of Transplantation*.

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