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COVID-19: A Global Transplant Perspective on Successfully Navigating a Pandemic

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ABBREVIATIONS:

ACE, angiotensin converting enzyme

BAL, bronchoalveolar lavage

COVID-19, Coronavirus disease 2019

HIV, human immunodeficiency Virus

ICU, intensive care unit

MELD, model for end-stage liver disease

MERS, middle east respiratory syndrome

NAT, nucleic acid testing

NP, nasopharyngeal

OPO, organ procurement organization

SARS-CoV, severe acute respiratory syndrome – coronavirus

TID, transplant infectious disease

Abstract

The COVID-19 pandemic has rapidly evolved and changed our way of life in an unprecedented manner. The emergence of COVID-19 has impacted transplantation worldwide. The impact has not been just restricted to issues pertaining to donors or recipients, but also health care resource utilization as the intensity of cases in certain jurisdictions exceeds available capacity. Here we provide a personal viewpoint representing different jurisdictions from around the world in order to outline the impact of the current COVID-19 pandemic on organ transplantation. Based on our collective experience, we discuss mitigation strategies such as donor screening, resource planning and a staged approach to transplant volume considerations as local resource issues demand. We also discuss issues related to transplant-related research during the pandemic, the role of transplant infectious diseases and the influence of transplant societies for education and disseminating current information.

Introduction

Transplantation has become an established treatment for end-stage organ diseases and is a highly regulated field. There are several threats to transplantation but one particularly important threat is that of an emerging infectious disease. Since the 1980s, there have been several emerging viral diseases including HIV in the late 1980s/early 1990s, SARS-CoV, West Nile Virus, pandemic influenza A/H1N1, Zika, Ebola, and now pandemic COVID-19 caused by SARS-CoV-2. For each of these threats, transplant programs have responded in a coordinated fashion by assessing the risk of donor transmission, assessing the severity of disease in the recipient, and recognizing the potential for transmission to healthcare workers¹⁻⁵. This knowledge has then been used to generate algorithms for donor screening, not using organs from potentially infected donors, and recipient management. Many of these emerging viruses have been manageable, sometimes only limited to certain geographic areas, and transplantation/donation has been able to adapt and continue to provide this life-saving therapy in a safe and effective manner. The current COVID-19 pandemic is unique and unprecedented in modern times. It has crossed borders and infected >180,000 persons worldwide that we know of, with likely many more undiagnosed cases. It has been difficult to contain partly due to the contagious nature of the virus and mild illness in a majority of individuals. Nevertheless, the emergence of COVID-19 has impacted transplantation worldwide. The impact has not been just restricted to issues around donors or recipients, but also health care resource utilization as the intensity of cases in certain jurisdictions exceeds available capacity. Based on our collective experience, we suggest mitigation strategies such as donor screening approaches, resource planning and a staged approach to transplant volume considerations as local resource issues demand. We also discuss issues related to the management of immunosuppression trials during the pandemic, and the role of transplant infectious diseases and transplant societies for education and disseminating current information. We believe our collective experience will be valuable to the transplant community in the absence of hard published research findings this early in the pandemic.

Approach to Donation

There is a potential for COVID-19 to be transmitted by organ donation although the risk of this is unclear and we are not aware of any reports of transmission. The virus is primarily isolated from the respiratory tract suggesting the lung is a very high-risk for transmission when used from an infected donor. However, virus is also been reported to be isolated from the blood in up to 15% of cases and therefore, all organs may be at risk of acquisition ⁶. With the SARS epidemic of 2003, autopsy data demonstrated virus in almost all organs including the liver, kidney, and intestines ⁷. Donor screening from both a clinical and laboratory perspective is therefore an important consideration and has been the subject of much discussion ⁸. In areas with significant community transmission, if organ donation is to proceed in a safe manner, the authors recommend that both clinical and rapid laboratory screening is required. This approach to donation may differ in countries depending on the degree of community-transmission of COVID-19. However, many areas have noted that due to limitations in test availability the true rate of community penetration may be unknown. During the SARS-CoV outbreak of 2003 in Toronto, a clinical donor screening tool was instituted, incorporating epidemiological and clinical features of the donor, which then allowed deceased-donor transplantation to continue ². However, unlike in 2003, there has been rapid development of nucleic acid testing (NAT) for SARS-CoV-2 and therefore, testing of nasopharyngeal specimens has been incorporated and is the cornerstone of donor screening algorithms in several jurisdictions. Real-time NP swab donor screening has been successfully deployed in organ procurement organizations (OPOs) within Canada, Italy, Spain, and South Korea. However, many questions remain, including the false negative rates of testing which can be due to inappropriate collection or a patient early in the incubation period. Since SARS-CoV-2 is known to use the ACE2 receptor for viral entry, a bronchoalveolar lavage (BAL) specimen may be more appropriate than naso/oropharyngeal swab. However, bronchoscopy would have the potential risk of aerosolization and may not be logistically feasible. For this approach to be successful, the test result must also be rapidly available. Laboratory-developed or commercial NAT testing needs to be made available to OPOs with results in hours. Ruling out COVID-19 in a donor is also essential for

the safety of organ procurement teams. Shortages of testing kits, reagents and laboratory resources to carry out donor screening in the midst of a pandemic is also a major consideration.

In Canada, we developed a COVID-19 donor clinical screening tool and also started NP swab NAT screening for COVID-19 (Figure 1). For areas with significant community-transmission, the tool could be modified to reduce the importance of travel history. In the latter case, NAT testing would play a much larger role but the tool allows a second layer of redundancy in the screening process. Collectively the authors have had experience with asymptomatic NAT positive donors.

In Switzerland, where virus is currently widely circulating in the community, we established universal screening for all deceased donors by NAT in NP swab or BAL on March 5, 2020 (Table 1). Given the potential increase in risk for health personnel and current limited resources in the ICU for performing bronchoscopy, we favor NP swab over BAL for screening of SARS-CoV-2. Of note, given the wide clinical presentation of COVID-19, potential donors can be asymptomatic or mildly symptomatic at the time of donation, thus highlighting the need for universal screening.

In Italy, where there is significant community transmission, deceased donor screening with NAT for SARS-CoV-2 on BAL has become mandatory starting from February 23, 2020. At the time of this manuscript none of the screened donors has been found positive. The main problem in Italy is the huge number of patients requiring mechanical ventilation and that many ICUs that have been transformed to COVID-19 ICUs. For this reason, the number of potential donors is expected to significantly decrease.

In Spain, with high community transmission, universal screening (through at least one NP specimen and, if feasible, one lower respiratory tract sample) is now mandatory for all lung and small bowel donors across the country. In addition, NAT screening is also required for any deceased donor with recent travel to or stay in selected high risk Spanish regions, contact with a confirmed COVID-19 case, with a positive symptom screen. No donor-derived transmission has been reported to date.

The Japan Society for Transplantation (JST) published their formal statement on March 6, 2020 (http://www.asas.or.jp/jst/pdf/info_20200306.pdf) and recommended to clinically screen donors for significant exposure to COVID-19, travel history to high risk countries, and symptoms including fever and respiratory symptoms. However, due to limited testing capacity, universal screening has not been adopted. Uniquely, JST recommended for both lung and liver living donors to stay at home or in hospital 14 days prior to avoid COVID-19 exposure, in cases where transplantation can be postponed for 14 days.

The Korean Society for Transplantation (KST) released their recommendation on March 13, 2020. The KST recommended that both living and deceased donors should be tested for SARS-CoV-2 NAT from NP swab prior to procurement. If the living donor and/or recipient visited the highly epidemic domestic regions (Daegu city or Gyeongsangbuk-do Province), or had any exposure, the transplant operation should be postponed for 14 days with close clinical monitoring.

Approach to Transplantation

In the face of a pandemic that may impact transplant recipients adversely, decisions to continue or cease transplantation need to be made by programs. Although donor transmission is an ominous possibility, many of these decisions are actually independent of donor transmission and have to do with the following considerations: 1) introducing immunosuppression into patients in the midst of a pandemic; 2) the risk vs benefit ratio of postponing transplant vs. proceeding; and 3) rationing of healthcare resources including both inpatient and outpatient resources. Lack of ventilator capacity is also an extremely important consideration once widespread activity is present. For example, kidney transplantation reduces the morbidity of dialysis and is cost-saving but is not immediately life-saving. The majority of cases of living donor kidney transplantation can be postponed without significant impact on the recipient. Although the clinical outcomes of transplant recipients with COVID-19 are not known, based on previous SARS and MERS publications, the mortality could be high and nasopharyngeal and tissue viral loads may be greater than in immune competence. Therefore, a decision must be made by individual programs whether

to newly immunosuppress patients and send them out into the community during the COVID-19 pandemic. The other part of the decision is how much healthcare resource is utilized by newly transplanted patients with regards to readmission rates and whether during a period of strained hospital resources, it would be appropriate to perform kidney transplantation. It has been suggested, that the risk of infection in the community for a newly transplanted recipient may be mitigated by either not using induction therapy or using an IL2 receptor antagonist for induction rather than polyclonal globulin induction. This is unknown but is a logical extension of data from other viral infections. Another possibility is to temporarily pause kidney transplantation but continue to transplant highly sensitized patients. This has been done in some jurisdictions.

The decision to pause life-saving transplants such as liver, heart and lung is more difficult as wait-list mortality is greater. However, for liver transplantation patients can be stratified based on MELD score and the decision made to only transplant high MELD patients.

Similarly, transplanting high status patients only may be a trade-off to completely stopping deceased donor transplantation.

We suggest a phased approach to decreasing transplant activity (Table 2). Such decisions need to be made collectively and depend on risk tolerance, hospital capacity and degree of virus activity in the jurisdiction. A phased approach can include a 25%, 50% and 75% trigger for activity reduction. Obviously in a situation where healthcare system is completely overwhelmed a 100% reduction may be unavoidable.

Another issue to consider is COVID-19 screening of recipients (clinical and/or laboratory) being admitted for transplantation. A standard screening form could be used. Whether asymptomatic patients should have a NP swab for COVID-19 NAT is debatable as this may be negative during the incubation period and may place unnecessary burden on resources.

Currently most of the authors' centers (with the exception of South Korea) are not screening asymptomatic recipients being admitted for transplantation with NAT.

Outpatient transplant clinics also need to be modified as hospitals prepare for increasing numbers of COVID-infected patients (Table 3). Elective transplant well-visits can be postponed and clinics can be scaled back such that only urgent visits need to be seen. If

resources are available, clinic staff may choose to screen all patients by telephone prior to the clinic visit for symptoms compatible with COVID-19. Telehealth/telephone calls can be substituted for such clinic visits. Transplant programs can direct patients to public health websites or transplant-specific websites for information.

Manpower issues are equally important. Members of the transplant team should not come to work if experiencing any symptoms compatible with COVID-19 and self-isolate if exposure has occurred. The authors note that rapid testing of transplant team members with even mild symptoms is very important. Transplant teams should have clear back-up plans should a team member become ill or quarantined. The authors centers have either disallowed vacation or allowed non-travel vacations with the understanding that they may be called in to help out – an ‘all hands on deck’ approach.

Although it is difficult to determine the course of the pandemic, the majority of hypotheses are based on peak of disease lasting several months followed by a stable level with either year-round or seasonal circulation of the virus. At some point transplant programs that have decreased activity will need to ramp up. We suggest this could be done in a phased approach where more urgent transplants could proceed first, with more ‘elective’ cases phased in later. This would need to be coupled with safe donation and transplant practices to prevent and treat COVID-19, as well as an understanding of local health resource limitations.

Role of Transplant Infectious Diseases

An increasing number of programs worldwide have transplant infectious disease (TID) specialists embedded in programs. During the COVID-19 pandemic, the TID physician can serve as a critical resource in an evolving environment. Although there are no publications on transplant recipients currently, transplant recipients have acquired COVID-19 and anecdotal information suggests that there is a spectrum of illness ranging from mild to severe disease. It is likely that these patients may also have higher viral loads with prolonged shedding. This may require longer quarantine periods until patients can be confirmed swab negative (e.g. we use 2 consecutive negative swabs to terminate

quarantine). With regards to treatment of an infected transplant recipient, the ID physician can be instrumental in obtaining and suggesting experimental therapies. Currently, there are many clinical trials and experimental therapies for COVID-19 including remdesivir, lopinavir/ritonavir, darunavir-cobicistat, interferon beta and (hydroxy)chloroquine as well as combinations of these therapies. Passive high titer immunoglobulin from recovered COVID patients has been used. Blunting the inflammatory response with a trial of corticosteroids is controversial^{9,10}. However, reduction of IL-6 in critically ill patients with tocilizumab has also been attempted. Detailed treatment algorithms are beyond the scope of this viewpoint, and are continually evolving as new data emerge. Numerous centers and groups have developed treatment guidelines for transplant and non-transplant patients. Antimicrobial management of the critically ill transplant patient (including potential drug interaction between antivirals and immunosuppressive drugs), an understanding of experimental therapies and clinical trials, as well as working with the transplant physician in reducing immunosuppression are all roles of a transplant infectious disease physician. The TID physician is also important at a programmatic level. Since surges in hospital capacity are predicted to occur with COVID-19, the TID physician can also determine whether non-COVID-19 patients with infections can be safely discharged home on oral or home IV therapies. Transplant ID physicians can also serve as an educational resource for other transplant physicians and coordinators, assist in answering patient queries regarding COVID-19. They can also be instrumental in building screening strategies for transplant patients, ie decide who can be safely managed as outpatient and who needs hospital admission. Overall, TID physicians can act as a linchpin of the transplant program during the COVID-19 pandemic by establishing safety procedures of donor and recipients, providing up-to-date educational tools, and prioritizing transplant activities according to evolution of the epidemiology of the pandemic.

Approach to Transplant Research Activities

An approach needs to be developed for centers that have ongoing clinical trials, especially those of novel immunosuppressives, in the face of a pandemic. To completely discontinue a trial and withdraw patients would generally not be required. However, modifications could be made such as halting enrollment of new patients. Patients already enrolled in a clinical trial should be advised to practice social distancing and frequent handwashing. Patients may not wish to come to hospital for study visits and this needs to be respected. Research teams should liaise with the sponsor for virtual visits. The hospital can consider a separate area for study bloodwork and administration or pickup of study drug where study outpatients are separated from patients that may be unwell.

On the other hand, research for COVID in transplantation is necessary and needs to continue. Some areas for research include outcomes in transplant patients, predictive diagnostics, and management strategies including optimal approach to immunosuppression adjustment. Vaccines are under development and will likely be in clinical trials. When this occurs a trial in transplant candidates and recipients is also warranted.

Safety of the Transplant Team

We must think about steps to take care of ourselves and ensure we remain healthy so that we can look after our patients. Risks to think about include: 1) teams traveling for donor procurement to areas of high risk, 2) performing high-risk procedures such as bronchoscopies on deceased donors, 3) team exposures to transplant recipients who may be shedding greater quantities of virus (so-called super-shedders or super-spreaders). Many of these risks can be mitigated by careful use of personal protection measures, and if needed avoidance of high-risk situations and following hospital directives regarding in-person meetings. Any transplant staff who is ill must immediately excuse themselves from work, and have appropriate testing as needed.

Organization of transplant programs with parallel teams (without contact among them) may be appropriate to continue with transplant activity even in case of transmission of cases

within health care workers. A very important but often neglected issue is the mental health of the transplant team. Anxiety and distress related to COVID-19 exposure concerns, transplant program closures and adverse outcomes in recipients all may contribute. Closely working with the transplant psychiatry team is a proposed strategy for addressing this.

Role of Transplant Societies

Transplant societies such as AST, ASTS, TTS, ISHLT and country-specific organizations have an important role as an educational resource in a rapidly evolving pandemic. Experts within societies should disseminate donor screening algorithms, share advice for transplant programs and management of transplant recipients. The American Society of Transplantation has rapidly created an information sheet for transplant professionals including suggestions for donor screening. In addition, information specific for transplant recipients has also been created. Similarly, The Transplantation Society has also provided information on COVID-19 for transplant professionals. Platforms such as webinars, virtual town halls, chat groups can be used to quickly exchange information. Transplant societies can also act as a conduit for research collaborations on COVID-19. For instance, the Spanish Group for the Study of Infection in Transplantation and the Immunocompromised Host (GESITRA-IC/SEIMC) has drafted a consensus guideline for the therapeutic management of transplant recipients with COVID-19 and has established a prospective multicenter registry to obtain real-time data from clinical experience.

Summary

This is an exceptional time for the world full of uncertainty and anxiety. For those of us working in transplantation, it is especially worrisome given the highly vulnerable group of patients we serve. In this context it is of the utmost importance that we come together as a team, to share knowledge and experience that will benefit all of our program and most importantly our patients.

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Data sharing is not applicable to this article as no new data were created or analyzed in this study.

Figure Legend:

Figure 1: COVID-19 Donor Screening Tool

Table 1: Summary of Donor and Recipient Screening Practices.

Country*	Deceased Donor Screening	Living Donor Screening	Pre-transplant Recipient Screening	Specimen Type
Canada**	Universal NAT	Universal NAT	Clinical	NP or BAL
Switzerland	Universal NAT	Universal NAT	Clinical	NP or BAL
Italy	Universal NAT	Universal NAT	Clinical	BAL in deceased donor NP in living donor
Spain	Universal NAT	Universal NAT; donation postponed 21 days if known exposure	Clinical	NP +/- BAL
Korea**	Universal NAT	Universal NAT	Universal NAT	NP
Japan**	Risk-based NAT due to limited testing capacity	Self-isolation or hospital-admission 14 days prior to surgery	Clinical; NAT where testing available	NP (and BAL for intubated patients)

NAT, nucleic acid testing, NP nasopharyngeal, BAL broncho-alveolar lavage

*may represent the author centers – not necessarily country wide; assumes transplant activity is continuing

**recommendations of country-specific transplant societies

Table 2: Phased Approach to New Transplant Activity During the COVID-19 Pandemic

Transplant Activity Level	Priority Level Description	Examples (may include but not limited to)
25% reduction in transplant activity	Elective cases. Patients whose conditions is deemed non-life threatening or can be managed with medication and for whom services can be deferred until the end of a pandemic wave (i.e. six to eight weeks.)	<ul style="list-style-type: none"> • Kidney Transplant <ul style="list-style-type: none"> ○ No living donor activity ○ Deceased donor activity allowed • Liver Transplant <ul style="list-style-type: none"> ○ No living donor activity for stable recipients ○ Deceased donor activity allowed • Heart – normal activity • Lung – normal activity • Kidney-Pancreas/Pancreas alone Transplant <ul style="list-style-type: none"> ○ K/P activity allowed ○ PAK or PTA not allowed • Islet Transplant <ul style="list-style-type: none"> ○ No Activity • Small Bowel <ul style="list-style-type: none"> ○ No Activity • Keratolimbal <ul style="list-style-type: none"> ○ No Activity
50% reduction in transplant activity)	Urgent cases. Patients who are deemed urgent and who need service within 14 days. It may be possible to defer these services for a few days, but not for the length of a pandemic wave. Physicians will determine that these patients are not put at undue risk. If their situation changes they will be changed to	<ul style="list-style-type: none"> • Kidney Transplant <ul style="list-style-type: none"> ○ No activity except highly sensitized (eg, PRA of 95% or above with suitable donor offer and negative DSA). • Liver Transplant <ul style="list-style-type: none"> ○ Activity for MELD > 25 ○ No living donor activity for stable recipients • Heart <ul style="list-style-type: none"> ○ Only intermediate status patients and above • Lung <ul style="list-style-type: none"> ○ All Patients allowed ○ Defer if patient is stable on wait list • Kidney-Pancreas Transplant <ul style="list-style-type: none"> ○ No Activity except high PRA as above • Small Bowel Transplant

	emergent	<ul style="list-style-type: none"> ○ No Activity ● Islet Transplant <ul style="list-style-type: none"> ○ No Activity ● No import of organs from select jurisdictions
75% reduction in transplant activity	Emergent cases: Patients who are deemed critical, whose condition is immediately life threatening. Their immediate need is greatest	<ul style="list-style-type: none"> ● Kidney Transplant <ul style="list-style-type: none"> ○ No activity unless for medically urgent status (eg, lack of dialysis access, uremic cardiomyopathy, uremic neuropathy with paralysis and/or respiratory compromise) ● Liver Transplant <ul style="list-style-type: none"> ○ Only Fulminant Hepatic Failure or MELD > 30 ● Heart Transplant <ul style="list-style-type: none"> ○ Highest Status Only (eg, Status 3, 3.5 and 4 in Canada) ● Lung Transplant <ul style="list-style-type: none"> ○ Only Rapidly deteriorating and status 2 patients (use LAS in U.S.) ● Kidney-Pancreas Transplant <ul style="list-style-type: none"> ○ No activity ● Small Bowel Transplant <ul style="list-style-type: none"> ○ No activity ● Islet Transplant <ul style="list-style-type: none"> ○ No Activity ● No import of organs from select jurisdictions
100% reduction in transplant activity	Health system is overwhelmed with COVID-19; No ICU or other capacity available; severe shortages of health personnel	<ul style="list-style-type: none"> ● Halt of all living and deceased donor transplant activity

PAK, pancreas after kidney PRA, panel reactive antibody PTA, pancreas transplant alone MELD, model for end-stage liver disease LAS, lung allocation score, ICU intensive care unit

Table 3: Ambulatory Transplant Clinic Service Reduction during COVID-19 Pandemic.

Transplant Activity Level	Transplant Centre Priority	Description (may include but not limited to)
25% Reduction in Transplant Activity	Should be Deferred	<ul style="list-style-type: none">• Annual & Well-visit Post-Transplant check ups• Consider telehealth• Non-urgent rehab• All blood work should be done at outside labs if possible• Redeploy Ambulatory Staff to establish 24/7 call center for patient
50% Reduction in Transplant Activity	Should be seen	<ul style="list-style-type: none">• Recent Transplant patients (Define for each organ)• Transplant Patients with sub-acute/chronic complications• Consider telehealth
75% Reduction in Transplant Activity	Need to be seen	<ul style="list-style-type: none">• Very Recent Transplants (Define for each organ)• Transplant Patients with acute complications to avoid inpatient admission
Near 100% Reduction in Ambulatory activity	No capacity due to health care system overwhelmed with COVID; lack of personnel;	<ul style="list-style-type: none">• Ambulatory activity unable to proceed. Referral to community primary care physicians if possible.

Figure 1: COVID-19 Donor Screening Tool*

SECTION A: Institution

This category describes whether institutional transmission of COVID-19 is of higher risk in the referring hospital. Unprotected COVID-19 exposure plus transmission from exposed person to others in donor ICU within the last 14 days: yes no

SECTION B: Active COVID-19 Infection

Has the potential donor been identified as a COVID-19 presumptive positive and/or as a COVID-19 confirmed case in the last 14 days? yes no

SECTION C: Exposure

Indicate whether the potential donor has had any of the following:

- Direct contact with known or suspected COVID-19 case in the last 28 days yes no
- Travel in the last 28 days to high risk region yes no
- Known COVID-19 diagnosis in the last 3 months yes no

SECTION D: Clinical

Indicate whether the potential donor has had the following signs or symptoms in the last 28 days, unless another explanation clearly exists.

- fever (> 38°C if taken) yes no
- felt unwell with myalgia and/or headache yes no
- persistent or frequent cough yes no
- shortness of breath yes no
- chest imaging showing lung infiltrates yes no

High Risk (NO DONATION and No Testing required):

ANY of

Section A: Yes answer

Section B: Yes answer

Moderate to High Risk: Testing required; Do not proceed without testing

Section C: Yes answer(s) with or without Section D: Yes answer(s)

Low to Moderate Risk: Testing required; Do not proceed without testing

Section D: Yes answer(s), with No answer(s) in Section A, B and C

Very Low Risk: Testing required; but MAY PROCEED WITH DONATION if test result cannot be obtained in time

No answer(s) in all of Section A, B, C and D

Footnote to Figure 1:

*Donor Screening Tool adapted from Trillium Gift of Life Network Organ Donation Organization, Ontario, Canada

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