STUDY PROTOCOL

Study Protocol for DeCOMPRESS: Defining the Disease Course and Immune Profile of COVID-19 in the Immunosuppressed Patient [version 1; peer review: 2 approved with reservations]

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\textbf{Abstract}

The ongoing coronavirus disease 2019 (COVID-19) pandemic is caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). Current advisory guidelines for high-risk groups—including people with autoimmune disease taking immunosuppressive therapies—are to take increased precautions and avoid any unnecessary contacts. The aim of the DeCOMPRESS study is to define the disease course and immune profile of COVID-19 in immunosuppressed patients. We will clinically phenotype patients with ANCA-associated vasculitis (AAV) who develop COVID-19 using a customized REDCap data collection instrument embedded within the Rare Kidney Disease (RKD) Biobank. This dataset will be interoperable with the rheum-COVID, Global Rheumatology Alliance, and SPRINT-SARI datasets, facilitating international data linkage. Acute and convalescent blood samples will be analysed by flow cytometry and ELISA to define the immunophenotype and cytokine profile. Patients will track COVID-19 and AAV symptoms through a bespoke smartphone app. DeCOMPRESS study findings will rapidly inform management of
immunosuppressed patients who contract COVID-19 by defining the natural history and immunological manifestations of the disease in these patients. We will also determine whether pre-existing immunosuppressant therapy lessens the cytokine storm associated with severe COVID-19 disease, thereby paradoxically improving rather than worsening clinical outcomes. This protocol document details the procedures for end-to-end completion of the DeCOMPRESS project and is complemented by an associated comprehensive Study Manual (accessible at: https://www.tcd.ie/medicine/thkc/decompress/).

Keywords
COVID-19, coronavirus, immunosuppression, autoimmunity, vasculitis, ANCA, cytokines, immunophenotype

This article is included in the Coronavirus (COVID-19) collection.
Introduction

Coronavirus disease 2019 (COVID-19) is a novel, infectious, multi-system disease caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). Clinical disease severity is highly variable, ranging from no symptoms, to a mild cough or breathlessness, to severe respiratory failure and death. The COVID-19 pandemic is particularly worrying for people with underlying health conditions, including those taking immunosuppressive medications for autoimmune diseases such as rheumatoid arthritis (RA), systemic lupus erythematosus (SLE), and anti-neutrophil cytoplasmic antibody (ANCA)-associated vasculitis (AAV).

The course of SARS-CoV-2 infection in patients with systemic autoimmune diseases who take immunosuppressive medication is poorly defined. Official guidance on strategies to prevent severe COVID-19 in patients taking immunosuppression is speculative, modelling of disease natural history in this population is ill-informed, and treatment for those who develop COVID-19 relies upon empirically reducing immunosuppression and providing “stress dose” glucocorticoids, without an evidence base to support this approach. Many older adults with autoimmune disease also have co-morbid conditions, including disease-associated organ damage, that might negatively impact their clinical outcomes.

However, early reports indicate that the most damaging facet of COVID-19 is a dysregulated “cytokine storm”, likely driven by dysfunctional neutrophils, suggesting that immunosuppression might confer more benefit than harm. Indeed, previous data suggest that biologic immunosuppressant agents might protect against severe sepsis (OR=0.56), and more recently glucocorticoids have shown efficacy as a treatment strategy for COVID-19.

Determining the risk vs. benefit posed by background immunosuppressive therapy, and devising strategies for optimal immunosuppressive management in the setting of COVID-19, has immediate clinical relevance. If immunosuppressive therapy is viewed as a vulnerability factor, rather than as risk-neutral or even protective, then a ward-based ceiling of care might erroneously be applied. Further, immunosuppressive therapy might be discontinued, risking a major disease relapse.

We will address this unmet need and contribute to the national and global COVID-19 response by analysing the impact of immunosuppression on the natural history of, and immune response to, SARS-CoV-2 infection. We will primarily use systemic vasculitis as a prototypical chronic autoimmune disease requiring long-term immunosuppression; however, findings will be broadly revealing about other chronic autoimmune diseases and will inform immunosuppressive medication management during this pandemic. The overarching goal of the DeCoPRESS study is to determine whether immunosuppressant therapy for chronic autoimmune disease protects against the cytokine storm associated with COVID-19 and reduces the severity of the clinical syndrome.

To achieve this goal, we will examine the following specific aims:

- Characterise immune cell subsets altered in COVID infection, comparing immunosuppressed to non-immunosuppressed persons;
- Profile serum cytokines in COVID infection, comparing immunosuppressed to non-immunosuppressed persons;
- Compare differences in clinical characteristics and outcomes between those who are versus those who are not immunosuppressed at the onset of COVID-19.

Study protocol

Study design

The DeCoPRESS study will use AAV as a chronic autoimmune disease model and take advantage of the existing Irish clinical research infrastructures built around this rare disease. This is a prospective multi-site cohort sub-study of the Rare Kidney Disease (RKD) Biobank, with collection of clinical data and biospecimens as defined in the primary RKD protocol, accessible at https://www.tcd.ie/medicine/thkc/research/rare.php. The RKD Data Management Plan and RKD Registry Data Dictionary are accessible at https://www.tcd.ie/medicine/thkc/decompress. Our key outcome measure is severe COVID-19 infection, defined as death, ICU admission and/or the need for high-flow (FiO2>30%) oxygen, non-invasive ventilation, or invasive ventilation.

Research Timeline

The start date for this project was June 1st, 2020 with an estimated timeframe of 24 months. Existing infrastructure supports the immediate implementation of this work.

Study Population – The RKD Biobank

The RKD Biobank cohort comprises over 850 members with systemic vasculitis. The three AAV phenotypes are microscopic polyangiitis (MPO), granulomatosis with polyangiitis (GPA), and eosinophilic granulomatosis with polyangiitis (EGPA). This bioresource, which includes about 70% of all known Irish AAV cases, has already supported many published studies. All recruits have provided biological samples during periods of remission and have consented to further sampling and data collection as envisaged for this project. All patients with systemic vasculitis enrolled in the RKD registry who develop probable or confirmed COVID-19 over the next 12–18 months will be included in the DeCoPRESS study.

Based on modelling data, we anticipate that approximately half of our enrolled patients will develop COVID-19 over the entire duration of the pandemic, and we will capture 75% of these cases (n=306). Based on empiric data (current medication usage in our cohort), we expect that two-thirds (n=202) will meet our definition of exposed (immunosuppressed) and one third (n=104) will not. We expect 52% of cases in the non-immunosuppressed group to develop severe infection. This expected sample size will have 80% power to detect an OR=0.5 (i.e. absolute frequency of severe infection of 35% in immunosuppressed group) or smaller, with 2-tailed alpha of 0.05.
Exposure groups. The patient group for this study comprises RKD Biobank recruits with a diagnosis of systemic vasculitis, as defined by the RKD protocol, and a diagnosis of COVID-19 as confirmed by polymerase chain reaction (PCR) at the time of diagnosis and/or by retrospective serologic testing. Patients with a high clinical suspicion of COVID-19 but no timely PCR test will be included pending confirmatory serologic testing. To estimate the overall period prevalence of SARS-CoV-2 infection in this population, including asymptomatic infection, all RKD recruits will be sampled between September 2020 and March 2021 (i.e., 6 to 12 months after the beginning of the pandemic) to determine the presence of SARS-CoV-2 antibodies. Review of the RKD database will be used to determine type, use, dosage, and frequency of immunosuppressive medications at the time of COVID-19 onset. Immunosuppressive exposure is defined as: current use of >5 mg daily prednisolone or any dose of azathioprine, mycophenolate, avacopan or methotrexate; or rituximab and/or cyclophosphamide use within the previous 6 months. The control group will comprise patients on minimal or no immunosuppression, defined as ≤5 mg daily prednisolone and no other immunosuppressant exposure, as defined above.

Study procedures
Patients with vasculitis previously recruited to the RKD Biobank will be identified, where possible, at the time of COVID-19 diagnosis. Patients will be treated at their local hospital or in the community and clinical data recorded in the RKD database. Blood samples will be obtained for immunophenotyping, serum cytokine measurements, and serologic analysis. This requires an efficient and responsive workflow to allow recruitment at short notice and outside office hours, something that the RKD infrastructure already allows for. However, due to delays in recognising cases or notifying study investigators, some patients might not be recruited until sometime after resolution of COVID-19; only convalescent immunophenotyping will be possible in these cases. A simplified workflow for the DeCOMpRESS study is described in Figure 1.

Sample and data collection
All participants in the DeCOMpRESS study will also be recruited to the RKD Biobank, either previously or at the time of COVID-19 diagnosis. Potential recruits presenting with COVID-19 at participating clinical sites will be reported in a de-identified manner as soon as possible to the DeCOMpRESS study team. There are two primary ways to identify suitable subjects for the DeCOMpRESS study:

1. Contemporaneously upon admission to hospital for COVID-19 treatment.
   a. Patients with vasculitis with a confirmed COVID-19 diagnosis (by PCR or serology) will be identified by their local medical team.
   b. Those who have not already been recruited to the RKD Biobank will be recruited to this before recruiting to DeCOMpRESS, as per RKD protocol (available at www.tcd.ie/medicine/thkc/research/rare.php).

2. Retrospective screening of local COVID-19 admission logs to identify RKD recruits, and of PCR/serology testing for SARS-CoV-2 in RKD recruits.
   a. The laboratory records of RKD recruits will periodically be screened for SARS-CoV-2 PCR testing, to identify any cases that may have been missed at the time of testing.
   b. SARS-CoV-2 serologic testing will also be periodically performed on routine RKD biobank study visits.

![Figure 1. A simplified workflow of the DeCOMpRESS study.](image-url)

Patients from designated cohorts are recruited at local hospital sites and clinical data recorded in REDCap. Serum and EDTA plasma samples are taken for immunophenotyping and serum cytokine analysis at the Central Pathology Laboratory in St. James’s Hospital. Repeat sampling at 3 and 6 months for convalescent antibody testing. Data will be incorporated into the ADAPT RDF data engine for subsequent analysis.
Upon identification of an eligible DeCoMPRESS study subject, the study team will follow the steps below:

1. Confirm RKD Main Study ID prior to the collection of clinical data or any study samples.
2. Notify the research team of a potential study subject, without disclosing any identifiable information.
3. Confirm an appropriate time for sample collection and transfer to a central biobank with the research team. This will ideally be less than 4 hours from sample collection to allow for efficient processing.
4. Collect 5 ml serum and 5 ml plasma samples as per RKD protocol.
5. Complete COVID-19 electronic case report form in RKD REDCap database.

Clinical data collection. Clinical data for the DeCoMPRESS study will be captured in an eCRF format using the REDCap platform. REDCap is a secure electronic data capture web platform that stores pseudonymised clinical data for the RKD biobank. The RKD registry database is hosted by Trinity College Dublin (TCD) Research-IT and access permissions are managed by the Principal Investigator (MAL). Comprehensive guidelines for completing the REDCap COVID-19 instrument are available in the RKD Clinical Data Entry standard operating procedure, and video tutorials for REDCap data entry are also available (accessible at www.tcd.ie/medicine/thkc/research/rare.php).

Registration fields are referred to as ‘instruments’ in REDCap. The detailed COVID-19 instrument records information about COVID-19 symptom onset, diagnosis, clinical features, management, and outcomes. Clinical data entry will be completed as soon as possible after patient recruitment to the study and no more than 7 days after biospecimen sampling. Particular attention will be paid to the COVID-19 instrument and prior and current vasculitis encounter medication entries. Patients will be classified as either exposed or unexposed based on immunosuppressant use.

Blood sample analysis
Immunophenotyping and serum cytokine analysis will be carried out at the central Pathology Laboratory (CPL) at St. James’ Hospital Dublin. Serum (minimum 5 ml) and plasma (minimum 5 ml) samples will be taken no more than 24 hours, and ideally within 4 hours, prior to analysis. Immune cell populations (percentages and counts of leukocytes) will be determined by flow cytometry. Serum will be archived to allow subsequent batch measurement of cytokines by ELISA and ELLA Multiplex cytokine assay. Cytokine analysis of historical (obtained during stable vasculitis remission before 1st September 2019), active infection, and convalescent samples will be performed, and compared both within and across participants, to determine the influence of immunosuppression, as well as COVID-19 disease severity, on cytokine levels before, during, and after COVID-19 infection. Additional details of immunophenotyping and serum cytokine analysis procedures are available in the DeCoMPRESS Study Manual (accessible at https://www.tcd.ie/medicine/thkc/decompress/).

Serology. All RKD subjects will be tested for antibodies (IgG, IgM and IgA) to SARS-CoV-2 -spike protein and receptor-binding domain of the spike protein using an in-house developed assay based on work from Mount Sinai19. The samples to be tested will be obtained:

1. 3 and 6 months post suspected or confirmed clinical infection.
2. From all RKD recruits (irrespective of clinical infection) between September 2020 and June 2021.
3. Before September 2019 (from archived serum in the RKD biobank).

A pilot series of patients with PCR-confirmed infection and 20 unselected RKD recruits with samples before and during COVID pandemic were tested for the purpose of this report. We will review patient-reported symptoms suggestive of possible COVID-19, logged via a customised smartphone app within the prior 12 months, and link this to the serological data. This will allow a comprehensive assessment of the impact of immunosuppressive therapy on the antibody response to SARS-CoV-2.

Integrated data analysis
Data analysis will employ the ADAPT Resource Description Framework (RDF) data integration engine and multivariable logistic regression with adjustment for demographics, comorbidities, and vasculitis disease activity11. Information from the multiple DeCoMPRESS data streams integrated into ADAPT’s analysis networks will be stored in a “triplestore” database to facilitate analysis and data sharing. This is a purpose-built database for the storage and retrieval of triples through semantic queries. Semantic triples codify statements about the data in the form of subject–predicate–object expressions. Data from various sources will be formatted appropriately to allow efficient uplift into the triplestore. The primary analysis outcomes will examine differences in immunophenotypes and cytokine profiles, clinical characteristics and outcomes between immunosuppressed and non-immunosuppressed patients. Additional exploratory analyses will include:

1. Differences in patient-reported symptom profiles (immunosuppressed vs. not)
2. Sub-group analyses comparing individual drug regimens e.g. induction vs. maintenance immunosuppression, rituximab vs. cyclophosphamide induction.
3. External comparisons to patients without vasculitis, with or without immunosuppression, who develop COVID-19.

Ethical approval and safety measures
DeCoMPRESS procedures are incorporated into the RKD ethical framework, with prior approval obtained in all RKD recruiting sites. All participants have provided consent for research projects deriving from the samples and clinical data. The
DeCoMPRESS sub-project has received additional approval by the RKD Biobank Steering Committee and from the SJH/ Tallaght Ethics Committee (REC: 2020-04 List 13 – Amendment (22)). The Health and Safety Authority has directed that university-based non-propagative work using SARS-CoV-2 positive samples should happen at containment level 2 using containment level 3 processes.12

Informed consent
All patients previously enrolled to the RKD Biobank have already provided informed consent for their data and specimens to be used in studies like DeCoMPRESS. Full details are provided in the RKD Biobank protocol. Patient information leaflets are available to download via the following link: https://www.tcd.ie/medicine/thkc/assets/pdf/RKD-Vasculitis-Patient-PIL-ICF-Version-5-07AUG19.pdf. In cases of COVID-19 in a patient with vasculitis who has not already been recruited to the RKD biobank, standard RKD recruitment procedures will be followed.

Statistical analysis
Descriptive analysis of group characteristics will include number and proportion of cases for categorical variables, mean and standard deviation (SD) for normally distributed continuous variables, and median and inter-quartile range (IQR) for non-normally distributed continuous variables. When examining associations between immunosuppression exposure and COVID-19 outcomes, statistical methods will be employed to account for confounding in vasculitis disease activity, e.g. multivariable logistic regression with handing of disease activity as a stratification variable or interaction term. Additional control groups, within and outside the RKD registry, may be identified and included in the study, pending appropriate ethical approval.

Dissemination
A key objective of DeCoMPRESS is rapid and effective dissemination of research outputs. This has been achieved primarily in conjunction with the Vasculitis Ireland Network (VINE)13, and UK and Ireland Vasculitis Society in conjunction with frequent reviews of state-of-the-art emerging research of relevance to the DeCoMPRESS mission. These are disseminated via a Twitter feed, targeted email distribution lists (managed by the UKIVAS secretariat) and using the VINE website. The DeCoMPRESS study is still recruiting patients and data will be compiled and published 18–24 after project commencement. Initial subject data has contributed to a manuscript with additional UKIVAS patient data and has been submitted for peer review. Full details of the wide-reaching dissemination strategy are outlined in the Data Sharing Plan (accessible at https://www.tcd.ie/medicine/thkc/decompress/).

Discussion
Current government advisory guidelines for patients taking immunosuppressive therapies encourage increased precautions to protect these vulnerable groups from contracting COVID-19. However, advice regarding how best to manage immunosuppressive therapies, both prior to and after developing COVID-19, are speculative. By comparing the disease course and immune profile of COVID-19 in patients with vasculitis who are versus are not immunosuppressed, the DeCoMPRESS study aims to determine the effects of immunosuppressant therapy on clinical outcomes in this patient group, which in turn will inform patient care. Based on these preliminary results it appears that in-hospital acquisition of COVID-19 during vasculitis investigation and management might be an important means of disease transmission in this population. The prevalence of COVID-19 in the vasculitis patient population appears broadly similar to the wider Irish population.

A key challenge of this study is our recruitment target. Based on epidemic modelling of COVID-19 incidence rates from April 2020, we predicted that approximately half of our cohort would develop COVID-19 over the course of the pandemic, of whom we targeted recruitment of 75% (n=306). However, since the design of this study, vulnerable patients, including those targeted in our study, have been advised to take additional precautions to prevent them from acquiring COVID-19. Further, serial lockdowns, and the emergence of an effective vaccine, might further (fortunately) impact recruitment to this study. Accordingly, in order to answer our research questions, we are exploring adding patients with other rheumatic and dermatological diseases, including those receiving immunosuppressive therapy as defined in this protocol. Further, we are planning more in-depth immunophenotyping to examine in detail the myeloid cell (neutrophil and monocyte) and innate T-cell response to SARS-CoV-2 that could be addressed with smaller patient numbers, should our original target not be reached.

Strengths of this study
Experience and diversity of study team. The research team were able immediately mobilize the appropriate resources to begin this study on the proposed start date of June 1st, 2020. Our core team and collaborators have a diverse expert knowledge in several areas of clinical and translational research, including immunology, epidemiology, autoimmune disease, and data analysis and integration. A patient representative from Vasculitis Awareness Ireland has also been included in our core research team to guide study design and will ensure that the best interests of patients are considered throughout the study, and that the research questions address issues of relevance to patients.

Access to patient resources via the RKD Biobank. The RKD Biobank comprises over 800 vasculitis patients, including about 70% of all prevalent Irish AAV cases. RKD Biobank recruits have already supported many studies14–17 and provided consent to use their clinical data and biological samples for research. The Vasculitis Ireland Network13 has been recognised by the Department of Health as a centre of expert vasculitis care. Data collection standards are comprehensive and consistent across all clinical study sites.

Powerful data infrastructures. Data generated throughout the DeCoMPRESS study will be uplifted into ADAPT’s RDF data integration engine.15 This uses semantic web technology to create a normalised data pool that can be queried and rapidly analysed as a single logical dataset. This will allow rapidly
harmonised dissemination of interoperable study data. We have generated a vasculitis-focused COVID-19 dataset, integrated into the RKD database, that adheres to FAIR data principles\(^{10}\). This was developed in collaboration with the FAIRVASC project, which seeks to merge multiple vasculitis registries across Europe, and is interoperable with international datasets. Finally, patients can monitor and report AAV and COVID-19 symptoms through a bespoke smartphone app, developed by Irish SME PatientMpower.

Data availability
No data are associated with this article.

References

1. Shahid Z, Kalayanamitra R, McClafferty B, et al.: COVID-19 and Older Adults: What We Know. J Am Geriatr Soc. 2020; 68(5): 926–929.
2. Gold MS, Sehayek D, Gabrielli S, et al.: COVID-19 and comorbidities: a systematic review and meta-analysis. Postgrad Med. 2020; 132(8): 749–755.
3. Pedersen SF, Ho YC: SARS-CoV-2: a storm is raging. J Clin Invest. 2020; 130(5): 2202–2205.
4. Hu B, Huang S, Yin L: Tissue damage from neutrophil-induced oxidative stress in COVID-19. Nat Rev Immunol. 2020; 20(9): 515–516.
5. Latorge M, Elbim C, Frère C, et al.: Tissue damage from neutrophil-induced oxidative stress in COVID-19. Nat Rev Immunol. 2020; 20(9): 515–516.
6. Schutte-Schrepping J, Reusch N, Pacilik D, et al.: Severe COVID-19 Is Marked by a Dysregulated Myeloid Cell Compartment. Cell. 2020; 182(6): 1419–1440.
7. Veras FP, Pontelli MC, Silva CM, et al.: SARS-CoV-2-triggered neutrophil extracellular traps mediate COVID-19 pathology. J Exp Med. 2020; 217(12): e20201129.
8. Richter A, Listing J, Schneider M, et al.: Impact of treatment with biologic DMARDs on the risk of sepsis or mortality after serious infection in patients with rheumatoid arthritis. Ann Rheum Dis. 2016; 75(9): 1667–73.
9. Ziegler CGK, Albon SJ, Nyquist SK, et al.: SARS-CoV-2 Receptor ACE2 Is an Interferon-Stimulated Gene in Human Airway Epithelial Cells and Is Detected in Specific Cell Subsets across Tissues. Cell. 2020; 181(5): 1016–1035 e19.
10. Amanat F, Stadlbauer D, Strohmeier S, et al.: A serological assay to detect SARS-CoV-2 seroconversion in humans. Nat Med. 2020; 26(7): 1033–1036.
11. Reddy BP, Houlding B, Hederman L, et al.: Data linkage in medical science using the resource description framework: the AVERT model [version 2; peer review: 2 approved]. HRB Open Res. 2018; 1: 20.
12. Laboratory biosecurity guidance related to coronavirus disease (COVID-19). [cited 2020 18/05/2020]. Reference Source
13. Trinity Health Kidney Centre: Vasculitis Ireland Network. [cited 2020 11/11/2020]. Reference Source
14. Dekkema GJ, Abdulahad WA, Bjima T, et al.: Urinary and serum soluble CD25 complements urinary soluble CD163 to detect active renal anti-neutrophil cytoplasmic autoantibody-associated vasculitis: a cohort study. Nephrol Dial Transplant. 2019; 34(2): 234–242.
15. Popat R, Hakki S, Thakker A, et al.: Anti-myeloperoxidase antibodies attenuate the monocyte response to LPS and shape macrophage development. JCI Insight. 2017; 2(2): e87379.
16. O’Reilly VP, Wong J, Kennedy C, et al.: Urinary Soluble CD163 in Active Renal Vasculitis. J Am Soc Nephrol. 2016; 27(9): 2966–16.
17. Pepper RJ, McAdoo SP, Moran SM, et al.: A novel glucocorticoid-free maintenance regimen for anti-neutrophil cytoplasm antibody-associated vasculitis. Rheumatology (Oxford). 2019; 58(2): 260–268.
18. Wilkinson MD, Dumontier M, Jan Albersberg IJ, et al.: The FAIR Guiding Principles for scientific data management and stewardship. Sci Data. 2016; 3: 160018.
Open Peer Review

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Silke Brix
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The ongoing coronavirus disease 2019 (COVID-19) pandemic is a global threat and a great challenge to patients with immune mediated inflammatory diseases requiring immunosuppression. The aim of the proposed study DeCOMPRESS is to define the disease course and immune profile of COVID-19 in immunosuppressed patients. It also aims to determine whether pre-existing immunosuppression lessens the cytokine storm associated with severe COVID-19 disease. This study could potentially assist in guiding the management of immunosuppressed patients contracting COVID-19 in the future.

The study rationale in respect to describing the COVID-19 disease course is sound with a sensible study protocol. The design is appropriate, but the key challenge will be the recruitment target. Due to the effective shielding programmes the numbers of infected vasculitis patients are not realistic within the study time frame. Collecting patients with different immune mediated inflammatory diseases will help to increase recruitment but will make the analyses more challenging due to significant varieties in disease processes and therapies. COVID-19 infections will however not fully subside in patients with immunosuppression. COVID-19 infections will likely continue to occur in immunocompromised patient cohorts due to the reduced vaccine response in these individuals and new viral variants. It might help to change the proposed timeline to achieve the study target. Given the vaccine role out, an important aspect will be COVID-19 disease behaviour in immunosuppressed patients after vaccination. I would suggest adding this aspect to the specific aims.

Regarding the immune profile: Given the severity of COVID-19 pneumonitis in certain risk groups, a fresh blood sample for immunophenotyping and serum cytokine analysis collected within 4hrs of diagnosis and prior to initiation of therapies such as steroids that significantly influence peripheral immune cells seems ambitious. A small sub-cohort seems more realistic and I would suggest aiming for 20-25 vasculitis patients with age-matched controls. The proposal measuring the serological long-term response to COVID-19 infection however is appropriate and a promising endeavour.
Is the rationale for, and objectives of, the study clearly described?
Yes

Is the study design appropriate for the research question?
Yes

Are sufficient details of the methods provided to allow replication by others?
Yes

Are the datasets clearly presented in a useable and accessible format?
Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: ANCA-associated vasculitides, biomarker of disease activity and clinical risk stratifications.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Author Response 23 Dec 2022

Emma Leacy, Trinity College Dublin, Dublin, Ireland

We thank the reviewer for their helpful comments. Given the dearth of evidence on the impact of COVID-19 in AAV patients, effective shielding programs was the primary protection measure in this cohort. Indeed there were very few infections noted in the first 9 months of this study. As vaccines became readily available we expanded the population to include sampling of healthy patients pre-and post-vaccination. The focus has always remained on AAV patients, but given the shared treatment strategies with other autoimmune conditions, our findings will be relevant to other patient groups. Further, the majority of samples were collected from the greater Dublin area and were indeed processed within 4 hours. This was made possible by the established infrastructure of the Rare Kidney Disease Biobank. We are aiming to publish full results of our vaccine cohort study in early 2023, and look forward to applying our findings to clinical practice.

Competing Interests: N/A

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I would like to commend the authors on undertaking this effort that would help address an unmet need in patients who are immunocompromised. The authors have the infrastructure and expertise to carry this ambitious work. I have a few suggestions/considerations for the authors:

1. The study's intended participants are patients with ANCA vasculitis. I am concerned that the information from this study might not be applicable to patients with other immune diseases due to several reasons. First, AAV frequently involves the upper and lower airways. Since COVID19 is a respiratory disease its behavior in patients with previous damage (due to AAV) may be very different from its behavior in patients with autoimmune diseases that do not have a high predilection for airways (such as SLE). Secondly, hypogammaglobulinemia is frequently encountered in patients treated for AAV but is not frequently observed in other autoimmune diseases (again such as SLE). This introduces a confounding factor that may render the findings of this study not generalizable to the rest of the immunocompromised population.

2. The authors mention that they will perform cytokine profiling. Adding more information about the cytokines of interest would be useful to the reader. If the authors are considering the use of multiplexed assays or a cytokine array would recommend stating the assays planned.

3. As vaccines become more available are the authors planning to gather vaccination data? It would be useful for comparison especially when it comes for key outcomes.

4. What secondary outcomes are the authors hoping to look into?

Is the rationale for, and objectives of, the study clearly described?
Yes

Is the study design appropriate for the research question?
Yes

Are sufficient details of the methods provided to allow replication by others?
Partly

Are the datasets clearly presented in a useable and accessible format?
Not applicable

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: ANCA vasculitis, lupus nephritis, glomerulonephritis

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.
We thank the Reviewer for their insightful comments, and have addressed them in order below.

1. Our primary patient population was patients with AAV. At the time of publishing this group had not been given sufficient attention in terms of infection nor vaccine response (see Strangfeld et al., 2020). Our group were in a unique position to be able to sample and study these patients, and built our study around them to be able to deliver specific, relevant information about how COVID-19 affects their disease. We did collect data on co-morbid conditions in our patient population, which will complement the results of other studies. Patients taking antibody-depleting therapies such as rituximab did suffer with hypogammaglobulinemia, and this is also a common treatment for other autoimmune conditions. Thus, this work is relevant to other immunocompromised populations. We will present findings of our comprehensive serology assessments in our results paper. Furthermore up to half of AAV patients (primarily PR3+) have pulmonary involvement (see Zhou et al., 2022, and Hunter et al., 2020), which further emphasises the need for specific, relevant advice on the impact of COVID-19 on their condition.

2. Cytokine measurements were carried out by multiplex ELLA assay (ProteinSimple) in the Clinical Pathology Laboratory in St. James' Hospital. IL-1β, IL-6, IL-8, TNF-α, and soluble CD25 were measured. Full details will be included in the Methods section of our results paper.

3. Data has been collected from patients before and after vaccination. Samples were collected from routine AAV clinics at Tallaght University Hospital prior to vaccination, and again 14-30 days after the second vaccination dose. Full details of sample collection and the associated results of this vaccination study are set to be published in 2023.

4. Our primary outcome was survival, and a secondary outcome measure was severity of infection graded as mild (did not require hospitalisation), moderate (required hospitalisation) or severe (required intensive care unit admission). Supplemental oxygen requirements were also recorded. In the vaccine response cohort our primary outcome measure was serology, and secondary measurements included changes in immune cell populations and serum cytokines.

**Competing Interests:** N/A