

DATE	
NAME	
GENDER	<input type="checkbox"/> Female <input type="checkbox"/> Male	
AGE	
BODY MASS	
HEIGHT	
ESRD underlying condition	<input type="checkbox"/> Diabetic kidney disease <input type="checkbox"/> Hypertensive nephropathy <input type="checkbox"/> Glomerulonephritis <input type="checkbox"/> Cystic/hereditary <input type="checkbox"/> Unknown <input type="checkbox"/> Other:	
Renal replacement therapy:	<input type="checkbox"/> Hemodialysis Start date:	
Concomitant diseases:	<input type="checkbox"/> Myocardial infarction <input type="checkbox"/> Congestive heart failure <input type="checkbox"/> Peripheral vascular disease <input type="checkbox"/> Cerebrovascular accident <input type="checkbox"/> Transient ischemic attack <input type="checkbox"/> Dementia <input type="checkbox"/> Chronic obstructive pulmonary disease <input type="checkbox"/> Connective tissue disease <input type="checkbox"/> Peptic ulcer disease <input type="checkbox"/> Diabetes mellitus <input type="checkbox"/> uncomplicated <input type="checkbox"/> end-organ damage <input type="checkbox"/> Hemiplegia <input type="checkbox"/> Chronic kidney disease	<input type="checkbox"/> Liver disease <input type="checkbox"/> mild = chronic hepatitis (or cirrhosis without portal hypertension) <input type="checkbox"/> moderate = cirrhosis and portal hypertension but no variceal bleeding history <input type="checkbox"/> severe = cirrhosis and portal hypertension with variceal bleeding history <input type="checkbox"/> Solid tumor <input type="checkbox"/> localized <input type="checkbox"/> metastatic <input type="checkbox"/> Leukemia <input type="checkbox"/> Lymphoma <input type="checkbox"/> AIDS <input type="checkbox"/> Other:
Kidney transplantation	Date: Donor <input type="checkbox"/> Living donor <input type="checkbox"/> Deceased donor Serum creatinine prior to COVID-19:	
Immunosuppressive regimen prior to COVID-19:	<input type="checkbox"/> Steroids <input type="checkbox"/> Tacrolimus <input type="checkbox"/> Cyclosporine A <input type="checkbox"/> MMF/MPA <input type="checkbox"/> Azathioprine <input type="checkbox"/> Sirolimus <input type="checkbox"/> Everolimus	
Immunosuppressive regimen modification after onset of SARS-CoV-2 infection:	<input type="checkbox"/> Dose reduction Drug name: Dose prior to reduction..... Dose after reduction..... <input type="checkbox"/> Drug withdrawal Drug name: <input type="checkbox"/> Other:	

SARS-CoV-2 vaccinations (brand name, dates):	<input type="checkbox"/> YES <input type="checkbox"/> 1 dose (date) brand name <input type="checkbox"/> 2 doses (date) brand name <input type="checkbox"/> 3 doses (date) brand name..... <input type="checkbox"/> 4 doses (date) brand name <input type="checkbox"/> NO	
SARS-CoV-2 vaccinations (side effects):	<div style="display: flex; justify-content: space-between;"> <div style="width: 48%;"> <input type="checkbox"/> Pain at the injection site dose number(s):..... <input type="checkbox"/> Fatigue dose number(s):..... <input type="checkbox"/> Headache dose number(s):..... <input type="checkbox"/> Muscle aches dose number(s):..... <input type="checkbox"/> Arthralgia dose number(s):..... <input type="checkbox"/> Chills dose number(s):..... </div> <div style="width: 48%;"> <input type="checkbox"/> Fever dose number(s):..... <input type="checkbox"/> Swelling at the injection site dose number(s):..... <input type="checkbox"/> Redness at the injection site dose number(s):..... <input type="checkbox"/> Nausea dose number(s):..... <input type="checkbox"/> Enlarged lymph nodes dose number(s):..... <input type="checkbox"/> Feeling unwell dose number(s):..... </div> </div>	
COVID-19:	date of the symptoms onset: symptoms duration (days): date of the positive test: hospitalization due to COVID-19 <input type="checkbox"/> YES <input type="checkbox"/> NO oxygen therapy <input type="checkbox"/> YES <input type="checkbox"/> NO minimal oxygen saturation (%).....	
COVID-19: symptoms	<input type="checkbox"/> anosmia <input type="checkbox"/> dyspnea <input type="checkbox"/> fever <input type="checkbox"/> cough <input type="checkbox"/> muscle and joint pain <input type="checkbox"/> weakness <input type="checkbox"/> diarrhea <input type="checkbox"/> dizziness <input type="checkbox"/> skin hyperalgesia <input type="checkbox"/> headaches <input type="checkbox"/> memory impairment <input type="checkbox"/> concentration problems	
MOLNUPIRAVIR THERAPY:	date of distribution: missed doses <input type="checkbox"/> YES <input type="checkbox"/> NO if yes, why premature therapy discontinuation: <input type="checkbox"/> WHEN: <input type="checkbox"/> WHY:	
MOLNUPIRAVIR THERAPY - side effects:	<div style="display: flex; justify-content: space-between;"> <div style="width: 48%;"> <input type="checkbox"/> diarrhea <input type="checkbox"/> nausea <input type="checkbox"/> vomiting <input type="checkbox"/> dizziness </div> <div style="width: 48%;"> <input type="checkbox"/> headaches <input type="checkbox"/> rash <input type="checkbox"/> other (please describe): </div> </div>	