# RKD Vasculitis COVID-19 Module

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The following was constructed as a means to capture data for immunosuppressed patients who contract COVID-19 (C-19). The aim was to ensure the module was granular, while still allowing for a busy clinician to complete it in a timely fashion. There is a large emphasis on interoperability (e.g. the use of standardised ontologies) and including fields to facilitate collaboration with other research groups. This C-19 module/instrument was developed using the following:

- SECURE-IBD (IBD C-19 registry) data dictionary
- COVID-19 Global Rheumatological Alliance data dictionary
- Data dictionary from https://rheum-covid.org/
- Health Protection Surveillance Centre (HPSC Ireland) Critical Care surveillance forms
- WHO COVID-19 surveillance CRF
- Comments from the UKIVAS group: Mark Little, Neil Basu, Rona Smith, Alan Salama, David Jayne, Silke Brix
- COVID-19 existing literature as of 23<sup>rd</sup> March 2020.

We have maintained existing relevant fields in existing instruments (visuals included below, and shaded in orange) and added an additional COVID-19 specific instrument, which was modelled on our current 'encounter' instrument (used for clinic visits) – the information will populate a simultaneous encounter (as a pop-up within the COVID-19 instrument) and other relevant fields (via piping).

#### **Dissemination:**

REDCAP has the capability of quick exports and rough raw statistics. The plan is to distribute a **weekly email** to give researchers and clinicians real-time information. R markdown is a potential option to standardise reports and graphics.

### Instrument (new): COVID-19

Field Label	Field Type	Choices
Reporter information		
Name of physician providing	Text	
care for vasculitis patient		
Name of centre providing care	Dropdown	Current hospital dropdown
		from 'Patient' – 'Hospital Unit',
		including 'other' as last option
Date this form completed	Calendar (Default = Today)	
Please select all databases this	Checkbox	Irish RKD / UKIVAS / EULAR /
form has been shared with		Global COVID

For 'Patient information', only non-orange fields appear in the COVID-19 instrument (as below):

Patient information (Part 1)		
Country of residence	B Ireland €	
County of residence	₩ →	
Education	₩	
E-cigarette or Vape use		

Patient information		
Gender	Dropdown	Male / Female / Other RKD: pipe in from 'baseline characteristics – common'
Race (select all that apply)	Checkbox	RKD: piping from 'ethnicity' in 'baseline characteristics – common'
Country of residence	Dropdown	RKD: Default = Ireland NEW RKD FIELD - below 'country of birth' in 'baseline characteristics – common'
County of residence	Dropdown	RKD: 32 counties listed in alphabetical order & 'other' with free text NEW RKD FIELD - below 'country of residence' in 'baseline characteristics – common'
Education	Dropdown	Primary School / High school / University / Unknown or Prefer not to answer <i>NEW RKD FIELD- above</i> <i>'employment status at</i> <i>diagnosis' in 'baseline</i> <i>characteristics – common'</i>
Smoking history	Dropdown	Current Smoker/ Former Smoker/ Never Smoker/ Unknown smoking status <i>RKD: pipe from smoking</i> <i>history in 'baseline</i> <i>characteristics – common'</i>
E-cigarette or Vape use	Dropaown	Yes / No / Unknown

		NEW RKD FIELD - add below the 'smoking history' field in 'baseline characteristics – common'
Comorbidities (check all that apply)	Checkbox	Asthma - SNOMEDCT/195967001 // Atrial fibrillation - SNOMEDCT/49436004 // Cerebrovascular accident (stroke) - SNOMEDCT/230690007 // COPD (not including late onset asthma) - SNOMEDCT/13645005 // Coronary heart disease (Ischaemic heart disease) - SNOMEDCT/414545008 // Diabetes mellitus - SNOMEDCT/73211009 // Hyperlipidaemia - SNOMEDCT/55822004 // Hypertension requiring treatment - SNOMEDCT/38341003 // Hypothyroidism - SNOMEDCT/40930008 // Renal disease (kidney disease) - SNOMEDCT/90708001 // Thromboembolic disease - SNOMEDCT/371039008 // Other (SNOMEDCT ontology) // None <b>RKD: pipe selected</b> comorbidities from 'pre- existing co-morbidities' instrument (see image below)

For **'Patient information (part 2)'**, all fields are obtained from existing fields, as explained for each variable. **It is important these fields are completed in the relevant sections** to ensure an enriched dataset:

Patient information (part 2)			
Vasculitis diagnosis	Dropdown	See standard UKIVAS options below	
		RKD: pipe in from 'Diagnosis – Vasculitis': either 'Small vessel	

		vasculitis (ANCA associated)'
		OR 'Small vessel vasculitis
		(Immune Complex)' OR
		'Medium vessel vasculitis' OR
		'Large vessel vasculitis' OR
		'Variable vessel vasculitis' OR
		'Single organ vasculitis'
ANCA serology (at time of	Dropdown	PR3 / MPO / PR3 and MPO /
diagnosis)		ELISA negative / No ELISA
		performed / Other (free text)
		RKD: pipe in from 'General
		characteristics – vasculitis' ->
		'At any point ANCA Specificity'
Date of formal vasculitis	Calendar	RKD: pipe in from 'Diagnosis –
diagnosis		Vasculitis': 'Date of diagnosis'
Biopsy performed	Dropdown	Yes / No / Unknown
		RKD: pipe in from 'Diagnosis –
		Vasculitis' – same field
Histologically confirmed	Dropdown	Yes / No / Unknown
diagnosis		
		RKD: pipe in from 'Diagnosis –
		Vasculitis' – same field
Vasculitis diagnosis confidence	Dropdown	Possible / Probable / Definite
Attach reference to diagnosis		RKD: pipe in from 'Diagnosis –
confidence table		Vasculitis' – same field

## For 'Vasculitis/Disease status at time of C-19 diagnosis':

'Encounters' Data: Vasculitis/Disease status at time of C-19 diagnosis BVAS Medication (partial)	
New Encounter Instance	
End-stage Kidney Disease (ESKD)	B ● Yes
VDI Score	₽

## • Complete ESKD and VDI Score, as detailed below:

End-stage Kidney Disease (ESKD)	Radio button	Yes / No RKD: piped to 'Renal Replacement Therapy' – 'End- stage kidney disease'
If ESKD = Yes Date of End-stage Kidney Disease (Date of	Calender	RKD: piped to 'Renal Replacement Therapy' – 'Date of End-stage kidney disease'

commencement on dialysis or transplant, whichever first)		
If ESKD = Yes		
Type of Renal Replacement Therapy (RRT)	Dropdown	Functioning renal transplant / Haemodialysis / Peritoneal Dialysis / Sustained CKD V
VDI Score	RKD: Integer	

- Then click 'New Encounter Instance', to reveal a pop-up window containing the usual 'encounter' instrument (see below). Complete this in full:
  - $\circ$  ~ Select 'yes' for 'COVID-19 related entry'
  - $\circ$   $\;$  Click 'Save & exit form' at the bottom of the pop-up after completing all fields
  - Then click 'x' in top right corner to close the pop-up

Rare Kidney Disease - RIT Pro	oduction - 03/09/19
- 11-15	
Current instance: 🔘 13 🗢	Data Access Group: TUH - Tallaght ?
Editing existing RKD ID 6 (Instance #13)	(Patient ld 6) 6
RKD ID	6
Patient ID	B View equation
Date Of Visit	H Today D-M-Y
* must provide value	COVID19 dataset, this value should be set to the same value as the 'Date this form completed' field on covid instrument
	○ Yes
COVID-19 related entry?	🖁 🔘 No
	COVID19 dataset
Interval from diagnosis (months)	Uiew equation

• This section includes the following variables:

Vasculitis/Disease status at time of C-19 diagnosis		
Employment status	Dropdown	Employed full-time / Employed part-time / Retired / Disability benefit / Student / Not working / Unknown
Disease activity at time of C-19 symptom onset or diagnosis if asymptomatic (physician global)	Dropdown	Active / Low disease activity / Remission
Urinalysis Done	Radio button	Yes / No
<i>If Urinalysis = Yes:</i> Urinalysis Protein	Radio button	Negative / +1 / +2 / > = +3
If Urinalysis = Yes:	Radio button	Negative / +1 / +2 / > = +3

Urinalysis Blood		
Last eGFR (CKD-EPI) prior to C- 19 diagnosis This field is obtained from the preceeding encounter for the bespoke export. No additional data entry is required for this	Integer	Field Note: Please enter NA if the patient was dialysis dependent at that time
Tield.	Padia huttan	Vac / Na
Darysis dependent		Field note: 'If patient has started/stopped dialysis since the last encounter, please input relevant dates into 'Renal Replacement Therapy' instrument.
If Dialysis dependent = yes	Calender	
Date of dialysis start +/- stop		
Weight (kg)	Number (allow 1 decimal	
	point)	
Height (m)	Please complete in 'baseline characteristics – common'	
BMI	Calculated field = weight (kg from above) / height <sup>2</sup> (m, from above)	
Clinical samples obtained	Tick boxes	<ol> <li>Plasma exchange fluid</li> <li>Serum</li> <li>Urine</li> <li>DNA extracted from saliva</li> <li>Saliva for DNA</li> <li>PAX gene tube for RNA</li> <li>EDTA for PBMC</li> <li>EDTA for DNA</li> <li>Tissue</li> <li>EDTA for plasma</li> <li>none</li> </ol>
BVAS Scoring	FULL BVAS (see SOP: clinical data entry for full explanation): radio button & tick boxes for each variable	
BVAS Score	Calculated field	
Do you think vasculitis is relapsing in this encounter? This is the physician's assessment at the time of	Dropdown	High probability / Possibly / No / Unknown
clinical review.		

Adjudicated probability of relapse?	Dropdown	Definite / High Probability / Possibly / No
This field is completed retrospectively by senior clinician, taking all variables into account (e.g. clinical signs/symptoms, laboratory values, biopsy, etc.)		Information link (hover over 'i in blue circle') to see definitions of Definite / High Probability, etc.

#### Medication

\*includes any medication the patient is on <u>within 2 weeks of COVID-19 diagnosis</u> AND \*all questions relate to drugs/doses at time of C-19 symptom onset (or diagnosis if asymptomatic)

Please ensure **'Treatment - continuing medication'** AND **'Treatment – intermittent pulse** administration' is up to date in terms of dose and start/stop dates for all immunosuppressive medications.

Immunosuppressive status	Dropdown	Currently on immunosuppression / Discontinuation of immunosuppression within 6 months prior to this encounter / Discontinuation of immunosuppression > 6 months prior to this encounter / Treatment Naïve
If Immunosuppressive status !=		
(not equals) 'Treatment Naïve'		
Corticosteroids	Radio button	Yes / No
If Corticosteroids = Yes		
Current corticosteroid dose	Radio button (RKD)	< 5  mg/day / 5-10  mg/day /
		$11_{20} \text{ mg/day} / 220 \text{ mg/day}$
		11-20 mg/ day / > 20 mg/ day
Corticosteroids in response to	Dropdown	Increased / No change /
this clinical encounter/enisode	Diopaoni	Reduced / Stopped / Unknown
If Immunosuppressive status !=		Select medication from
(not equals) 'Treatment Naïve'		dropdown list. If it is not listed.
(		select 'other' and enter the
Immunosuppressive	Drandown	medication in the new field
madiantian	Diopadwin	that appears using an ATC
		had appears, using an ATC
		backed ontology (i.e. when
For implementation:		you begin to type the ontology
		will suggest medications to

Patients may be on >1		select).
Immunosuppressive agent. If		
so, select the second agent		The following are a list of
from the field 'Additional		medications that we are
Immunosuppressive		particularly interesting in
medication. Consequently,		capturing (with the associated
complete whether this		ACI ONTOIOGY):
additional agent was		Alfa1 antitrypsin – UATC/B02AB02
Increased / no change /		Anakinra – UATC/L04AC03
reduce, etc.		Apremilast – UATC/L04AA32 Azathioprine - UATC/L04AX01
If the patient is not on any		Belimumab – UATC/L04AA26
additional agents, leave this		Bevacizumab – UATC/L01XC07
blank.		Ciclosporin – UATC/L04AD01
		Chloroquine – UATC/P01BA01
		Cyclophosphamide (Daily Oral) –
		Cyclophosphamide (Intravenous pulse) –
		UATC/LC01AA01
		Hydroxychioroquine – UATC/P01BA02 IVIG: Immunoglobulins – UATC/J06BA02
		Leflunomide - UATC/L04AA13
		Mepolizumab - UATC/R03DX09
		Mycophenolate mofetil - UATC/L04AA06
		Plasma (from recovered patients) –
		UATC/B05AX03 // Rituximab - UATC/L01XCO2
		Secukinumab – UATC/L01AC10
		Sulfasalazine – UATC/A07EC01
		Tacrolimus (including Advagraf, Prograf, etc.) - UATC/104AD02
		Thalidomide – UATC/L04AX02
		Tocilizumab – UATC/L04AC07
		Tumor necrosis factor alpha (TNF-)
		inhibitors – UATC/L04AB
		Ustekinumab - UATC/L04AC05
		None // Unknown
Immunosuppressive	Dropdown	Increased / No change /
medication in response to this		Reduced / Stopped / Unknown
clinical encounter/episode		
If Immunosuppressive	Dropdown	Clear response (i.e. improved
medication in response to this		clinically) / No response /
clinical encounter/episode =		Unknown response
Increased:		
Response to increased		
immunosuppresion		

- The remaining encounter instrument includes fields surrounding prophylactic medication and laboratory results. Please see the SOP: clinical data entry for further instructions.
- For Investigations summary: XXX
- Other medication variables to complete on the main C19 instrument include:

Angiotensin-converting-	Radio button	Yes / No / Unknown
enzyme inhibitor at C19		
diagnosis (ACE-i)		
Angiotensin II receptor blocker	Radio button	Yes / No / Unknown
at C19 diagnosis (ARB)		
Non-steroidal anti-	Radio button	Yes / No / Unknown
inflammatory drug at C19		
diagnosis (NSAID)		

COVID-19 Questions		
Date of C-19 symptom	Calendar	
onset (if known)		
Date of C-19 diagnosis	Calendar	
Interval (days) between	Calculated field	
symptom onset and	using 'date of C19	
diagnosis (if known)	symptom onset'	
	and 'date of C19	
	diagnosis'	
Age at C-19 diagnosis	Calculated field	
(years)	using DOB	
	('Baseline	
	characteristics –	
	common')	
Location at which C-19	Dropdown	1, Home or standalone testing   2, Nursing
diagnosis was made		home or assisted living facility   3,
		Outpatient facility   4, Emergency Room   5,
		Inpatient/Hospital   6, Other (freetext)   7,
		Unknown
Method of C-19 testing	Dropdown	1, symptoms (presumptive)   2, PCR   3,
(select the most objective		antibody   4, metagenomic testing   5, CT
option)		scan   6, other (free text)  7, Laboratory
		assay, type unknown
If Method = PCR:		
Level of Sars-CoV-2 (COVID-	Integer	
19) PCR		
Sars-CoV-2 (COVID-19) IgM	Integer	
Sars-CoV-2 (COVID-19) IgG	Integer	
Admission to hospital	Radio button	Yes /No/ Unknown
required		
If Yes:	Calendar	
Date of admission		
Admission to Intensive Care	Radio button	Yes /No/ Unknown
Unit		
If Yes:		
Date of admission	Calendar	
Interval (months) from AAV	Calculated field	
diagnosis to C-19 diagnosis	using Date of	

	diagnosis ('Diagnosis – vasculitis')	
Have patient's symptoms resolved at time of this report?	Dropdown	1, yes   2, no   3, unknown   4, Asymptomatic patient (just tested positive)
If Have patient's symptoms resolved at time of this report? = yes		
Date of symptom resolution (if known)	Calendar	Field note: first date patient is asymptomatic, signifying recovery
Interval (days) between symptom onset (if known) and symptom resolution OR current date (if symptoms persist)	Calculated field using 'date of symptom onset' and 'date of symptom resolution' (if not known use 'today's date')	
Date of hospital discharge (if known)	Calendar	
Length of stay (days)	Calculated field using 'date of admission' and 'date of hospital discharge'	
Infection Acquisition	Dropdown	High-risk travel to endemic area / Contact of known or suspected person / Attendance to a healthcare facility/ward where C-19 infections are managed / None of the above (community acquired) / Unknown
Clinical features at outset (check all that apply)	Checkbox	Fever – MEDDRA/10016558 / Malaise – MEDDRA/10025482/ Headache – MEDDRA/10019211/ Irritability or confusion – MEDDRA/10010300 / Arthralgia – MEDDRA/10003239 / Myalgia – MEDDRA/10028411 / Conjunctivitis – MEDDRA/10010741 / Rhinorrhea – MEDDRA/10039101 / Anosmia – MEDDRA/1002653 (Loss of smell) / Ageusia – MEDDRA/1001480 (Loss of taste) Sore Throat – MEDDRA/10041367 / Cough – MEDDRA/10011224 / Sputum production – MEDDRA/10041805 / Shortness of Breath – MEDDRA/10040604 / Chest pain – MEDDRA/10008479 / RR >24 breaths/min – MEDDRA/10038711/

		Abdominal pain – MEDDRA/10000081/ Nausea – MEDDRA/10028813 / Vomiting – MEDDRA/10047700 / Diarrhea – MEDDRA/10012735 / Other (MEDDRA Ontology) /
		None (asymptomatic) / Unknown
Body temperature ( <u>highest</u> <u>recorded</u> , °C)	Number (allow 1 decimal)	
	Validation: only allow range 13 - 46°C	
CRP (mg/L) Field note: Please enter the	Number (allow 1 decimal)	Some lab values are entered in the encounter pop-up (as described in the sop: clinical data entry) and the remaining C19
highest/peak recorded lab value for each field	RKD: for all lab values pipe to corresponding field in 'encounters'	specific lab values are completed in the COVID-19 instrument (these are highlighted in yellow).
Creatinine (mmol/L)	Integer	
AST (U/L)	Integer	
ALT (U/L)	Integer	
Haemoglobin (g/dL)		
Total White Cell Count x 10 <sup>9</sup> /L	Number (allow 1 decimal)	
Neutrophil count x 10 <sup>9</sup> /L	Number (allow 1 decimal)	
Lymphocyte count x 10 <sup>9</sup> /L	Number (allow 1 decimal)	
Neutrophil / Lymphocyte ratio	Calculated field	
Urine Protein Creatinine ratio (uPCR, mg/mmol)		
ANCA IF	Dropdown	Atypical / C / P / Negative / Not tested / Pending
Anti-PR3 level	Integer	
Anti-MPO level	Integer	
Platelet count x 10 <sup>9</sup> /L	Integer	
Creatinine kinase (U/L)	Integer	
D-dimer (mg/L)	Integer	
Ferritin	Integer	
Lactate (mg/dL)		
Prothrombin time (s)	Number (allow 1 decimal)	
Lactate dehydrogenase (U/L)	Integer	
Troponin	Integer	
Next to 'Troponin': Units	Dropdown	

Findings on chest imaging	Dropdown	Inflammation Ischaemia Infarction Haemorrhage Bony destruction Metastasis Mass/tumour Abscess Polyp Ulceration Cirrhosis Fibrosis Calcification Effusion Effusion Effusion Effusion Effusion Embolism Nodules Consolidation Organomegaly Single granuloma Multiple granulomas None
Were antibiotics administered?	Radio button	Yes / No / Unknown
Was treatment administered for C-19 infection (other than best supportive care)?	Checkbox	No treatment except supportive care Kaletra ((Lopinavir/ritonavir) – UATC/J05AR10 Remdesivir – <b>No ontology code yet as novel</b> Chloroquine – UATC/P01BA01 Hydroxychloroquine – UATC/P01BA02 <b>Neuraminidase inhibitors, direct acting antivirals</b> (e.g. Oseltamivir) – UATC/J05AH Azithromycin – UATC/S01AA26 Tocilizumab – UATC/L04AC07 Bevacizumab – UATC/L04AC07 Bevacizumab – UATC/L04A29 Alfa1 antitrypsin – UATC/B02AB02 Ciclesonide – UATC/R03BA08 Plasma (from recovered patients) – UATC/B05AX03 Other – ATC backed ontology
Complications / Disease Course (check all that apply)	Checkbox	Acute Respiratory Distress Syndrome – MEDDRA/10001052 / Acute respiratory failure (Type 1/2) – MEDDRA/10001053 / Pneumothorax – MEDDRA/10035759 / Acute liver injury – MEDDRA/10067970 / Acute heart failure – MEDDRA/1000803 / Myocarditis – MEDDRA/10028606 / Cardiac arrhythmia – MEDDRA/10003119 / Cardiac ischaemia – MEDDRA/10007584 / Acute Kidney Injury (AKI) – MEDDRA/10069339 / Sepsis – MEDDRA/10040047 / Vasopressor dependence at any time – MEDDRA/10064148 / Disseminated Intravascular Coagulation – MEDDRA/10013442 / Severe anaemia – MEDDRA/10002082 / Gastrointestinal haemorrhage – MEDDRA/10017956 /

		Encephalitis – MEDDRA/10014581 / Pregnancy-related complications – MEDDRA/10036569 / Hyperglycaemia – MEDDRA/10020637 / Hypoglycaemia – MEDDRA/10020996 / Rhabdomyolysis – MEDDRA/10039020 / Metabolic acidosis – MEDDRA/10027417 / Secondary infection – MEDDRA/10062158 / Macrophage activation syndrome – MEDDRA/10053867 / Other (MEDDRA ontology) / None
If AKI = selected, then please complete the		
following fields in the encounter pop-up.		
Dialysis dependent, Date of dialysis start +/- Date of dialysis stop		
Concomitant respiratory pathogens detected (select all that apply):	Checkbox	Influenza A Influenza B NON-COVID-19 Coronavirus Respiratory syncytial virus (RSV) Adenovirus
If Secondary Infection = selected Secondary infection is selected, please indicate the type of Infection (in addition concomitant respiratory viral pathogens)	Dropdown	Bacteremia (5758002) Pneumonia (233604007) Pyelonephritis (4581600) Gastroenteritis (25374005) Encephalitis (45170000) Cellulitis (128045006) Osteomyelitis (60168000) / Other Other using SNOMEDCT ontology
If Secondary Infection = selected		
With regards to the secondary infection, please select the microorganism (if known)	SNOMEDCT ontology	Begin typing the name of the microorganism into the field and the ontology will begin to suggest matches.
C-19 Outcome (Select the highest level of support the patient received)	Dropdown	<ol> <li>Not hospitalized, no limitations on activities</li> <li>Not hospitalized, limitation on activities</li> <li>Hospitalized, not requiring supplemental oxygen</li> <li>Hospitalized, requiring supplemental oxygen</li> </ol>

		5. Hospitalized, on non-invasive ventilation
		or high flow oxygen devices
		6. Hospitalized, on invasive mechanical
		ventilation or ECMO
		7. Death
		8. Unknown
If C-19 outcome = 7. Death		
-		
Date of death	Calendar	
	DKD, pipe to (date	
	RKD: pipe to date	
	of event in	
	baseline	
	characteristics –	
	common'	
If C-19 outcome = 7. Death		
Cause of death	SNOMEDCT	
	ontology	
	RKD: pipe to 'cause	
	of death' in	
	'baseline	
	characteristics –	
	common'	
May we contact you to get	Radio button	Yes / No
more information about the		
outcomes of this case?		
Would you like to share any	Text	
lessons or other aspects		
from this case? Please		
include as much		
information as desired this		
will greatly help natients		
and colleagues		
and colleagues.		

## Current relevant instruments in RKD Registry

Relevant fields identified with red box. For more details please refer to SOP: Clinical Data Entry

Baseline characteristics - common	
Editing existing RKD ID 10 (Patient Id 10) 10	
RKD ID	10
Gender	Eremale
Date of Birth	H D-M-Y
Year of birth	H 1985 View equation
Ethnicity	(H) → W2 - White Irish
Consent obtained for registry/biobank	<ul> <li>Consent for use of data in registry</li> <li>Biobanking</li> <li>Future contact</li> <li>All</li> <li>No</li> </ul>
Consent Version Number	H
Date of Initial Consent	(H) 21-12-2012 第 Today D-M-Y
Has consent been obtained for Genetic studies	H 🗘
Employment status at diagnosis	B Not working ♦
Ethnicity of mother	B NS - Not Stated ♦
Ethnicity of father	B NS - Not Stated ♦
Country of birth	(H) ↓ (Ireland ◆
Relationship to other case	H +
Maternal or Paternal	H • •
Clinical samples obtained	<ul> <li>Plasma exchange fluid</li> <li>Serum</li> <li>Urine</li> <li>DNA extracted from saliva</li> <li>Saliva for DNA</li> <li>PAX gene tube for RNA</li> <li>EDTA for PBMC</li> <li>EDTA for PDNA</li> <li>Tissue</li> <li>EDTA for plasma</li> <li>none</li> </ul>
Name of biobank biological sample is stored in	
Height	⊖ Unknown €

Smoking History	
Smoking	Previous 🕈
Age of starting	<ul> <li>(H)</li> <li>(P)</li> <li>(P)</li></ul>
Date of stopping	H 18-03-2020 Today D-M-Y
Average no. of cigarettes per day	H 20
Patient Status	
Status	B Dead ♦
Death	<ul> <li>IF ○ Yes</li> <li>IF ○ No</li> </ul>
Date of event	H Today D-M-Y
Cause of death	
Cause of death - Known	
Cause of death - known, snomedCT	Coronavirus infection 186747009
Torm Status	
Complete?	Complete

## 📱 Renal Replacement Therapy

Editing existing RKD ID 10 (Patient Id 10) 10	
RKD ID	10
Required renal replacement therapy during first presentation	<ul> <li>⊢ ● Yes</li> <li>⇒ ● No</li> </ul>
<b>Renal recovery</b> (independence from dialysis, regardless of dialysis duration)	<ul> <li>⊢ O Yes</li> <li>⇒ O No</li> </ul>
End-stage kidney disease	<ul> <li>Yes</li> <li>No</li> <li>reset</li> <li>Dialysis for &gt;90 days, sustained CKD V, commencement of dialysis and death within 90 days, and/or transplantation</li> </ul>
<b>Date of end-stage kidney disease</b> (date of commencement on dialysis or transplant, whichever first)	H 18-03-2020 Today D-M-Y If patient develops ESKD after initially recovering from dialysis- dependent kidney failure, enter the date that ESKD develops
Form Status	
Complete?	H Incomplete ♦
	Save & Exit Form Save & Stay Cancel

Editing existing RKD ID 10 (Patient Id 10) 10	
RKD ID	10
Please enter only one comorbid event per instance.	
Comorbid events (present before vasculitis diagnosis)	<ul> <li>Asthma - SNOMEDCT/195967001</li> <li>Atrial fibrillation - SNOMEDCT/49436004</li> <li>Cerebrovascular accident (stroke) - SNOMEDCT/230690007</li> <li>COPD (not including late onset asthma) - SNOMEDCT/13645005</li> <li>Coronary heart disease (Ischaemic heart disease) - SNOMEDCT/414545008</li> <li>Diabetes mellitus - SNOMEDCT/73211009</li> <li>Hyperlipidaemia - SNOMEDCT/55822004</li> <li>Hypertension requiring treatment - SNOMEDCT/38341003</li> <li>Hypothyroidism - SNOMEDCT/40930008</li> <li>Malignancy (neoplasm) - SNOMEDCT/108369006</li> <li>Renal disease (kidney disease) - SNOMEDCT/9708001</li> <li>Thromboembolic disease - SNOMEDCT/371039008</li> <li>Other</li> <li>None</li> </ul>
Comorbid events (present before vasculitis diagnosis)	
Comorbid events detail	
Form Status	
Complete?	⊖ Complete ♦

🗧 Diagnosis - vasculiti	sis - Vasculiti	Diagnos
-------------------------	-----------------	---------

📱 Diagnosis - Vasculitis	
Editing existing RKD ID 10 (Patient Id 10) 10	
RKD ID	10
Date of diagnosis	H Today D-M-Y
Age at diagnosis	H View equation
Date of onset of symptoms - known/unknown	🛞 Unknown 🗘
Date of onset of symptons	H Today D-M-Y
Age at onset	View equation
Small vessel vasculitis (ANCA associated)	$\bigcirc$ Granulomatosis with polyangiitis (Wegener) - ( )
Please select up to a MAXIMUM of 3 diagnoses	
Small vessel vasculitis (Immune complex)	₩ →
Medium vessel vasculitis	
Large vessel vasculitis	
Variable vessel vasculitis	₩ →
Single organ vasculitis	
Secondary vasculitis	<ul> <li>⊢) ● Yes</li> <li>◯ No</li> </ul>
Other	<ul> <li>(i) ● Yes</li> <li>(ii) ● No</li> </ul>
Other details	
Unclassified	<ul> <li>⊢ ○ Yes</li> <li>○ No</li> </ul>
Biopsy performed	H Yes
Histologically confirmed diagnosis	H Yes 🗘
Please open supporting PDF for reference	
Diagnosis confidence (initial assessment)	B Definite
Form Status	
	(H)

Treatment – continuing medications (PO)

Drug	Dose	Unit of Dose	Start Date	Stop Date
UATC/H02AB06	5	mg	2013-02-03	2014-10-16
2 Prednisolone - UATC/H02AB06	10	mg	2014-10-16	2018-01-05
3 Prednisolone - UATC/H02AB06	5	mg	2016-07-22	2017-12-10
1 Other	40	g	2016-11-08	
urrent instance: 🔘 g	5 🗢			
🥜 Editing existing RKD	DID 10 (Instance	<b>#5)</b> (Patient ld <b>10</b> ) 10		
RKD ID			10	
Drug				
Drug, DRON				
			(H)	
Drug			Other	÷
Drug, ATC			H raminril	C094405
				007000
Dose			H 9 5	
			(f)	
Unit of Doses			o (mg	<b>\$</b>
Frequency			Baily	<b></b>
,				•
Start Date			8-03-2020	Today D-M-Y
On going			🕒 🖲 Yes	
Stop Date				Today D-M-Y
Form Status				
Commission 2			θ	
Complete?			incomplete	

## Treatment – intermittent pulse administration (IV)

Editing existing RKD ID 10 (Patient Id 10) 10					
RKD ID					
IV therapy	Cyclophosphamide Injectable Solution - UATC/ L01AA01 Rituximab - UATC/L01XC02 Mabthera				
Date of IV therapy	Rituximab - UATC/L01XC02 Truxima Methylprednisolone - UATC/D07AA01 Ivig - UATC/J06BA02 (immunoglobulins, normal huma	Rituximab - UATC/L01XC02 Truxima Methylprednisolone - UATC/D07AA01 IvIg - UATC/J06BA02 (immunoglobulins, normal human, for intravenous administration)			
Dose of IV therapy	Mepolizumab - UATC/R03DX09 Other				
Unit of dose	😬 (mg 🔶				
Form Status					
Complete?	😁 Complete 🗘				

#### References

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Huang C, Wang Y, Li X, et al. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. Lancet 2020;395:497–506.

Xu X-W, Wu X-X, Jiang X-G, et al. Clinical findings in a group of patients infected with the 2019 novel coronavirus (SARS-Cov-2) outside of Wuhan, China: retrospective case series. BMJ 2020;368:m606.

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Metlay JP, Waterer GW, Long AC, et al. Diagnosis and Treatment of Adults with Community-acquired Pneumonia. An Official Clinical Practice Guideline of the American Thoracic Society and Infectious Diseases Society of America. Am J Respir Crit Care Med 2019;200:e45–67.

HPSC Surveillance:

https://www.hpsc.ie/az/respiratory/coronavirus/novelcoronavirus/surveillance/ADULT%20influenza%20&%20COVID%201 9%20ICU%20Admission&%20Discharge%20form 20200303 %20v032.pdf