

DIAMONDS CRF - patients with suspected COVID19

PATIENT DETAILS (this box should NOT be transmitted outside local hospital):									
Surname:	Hospital or NHS number:								
First name:	Search patient identification log to see if the patient already consented to the study. If yes, use the previously assigned study number and a new episode number								
Partial postcode (UK): - X X	assigned stady number and a new episode number								

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Study	/ number:		-1		- E	
Jiuu	/ IIuIIIDEI .					

DIAMONDS CRF – patients with suspected COVID19

1.1 STUDY ID DET	TAILS (DIS-C	entre ID-subje	ect ID-episode; e	g DIS-1101-1001-E01)	
Site code (eg 1101):	_ _ [OOB (ddmmyy):		
Study number			∕lale □	Female	
Episode (eg 01)		A	Admission heigh	nt (cm) Weigh	t (kg)
Alternate study nu	ımber(s)				
Consented to use re	esearch samples Yes [No 🗌 🤇	Consented to sha	re samples? Yes [□ No □
1.2 ADMISSION T	IMELINES				
	Date of	symptom ons	et D D N	M Y Y	
	Dat	e of fever ons	et D D N		
Date & t	ime of first presentation	to any hospit	al D D N	VI W	H H M M
	A	dmitted to ITU	? Yes ☐ N	No 🗌	
	Date & time	admitted to IT	U D D N	И И У	H H M M
	Date & time of first r	esearch blood	ls D D N	И И У У	H H M M
		Consent da		M M Y Y	
1.3 PRESENTING I			VITAL SIGNS AT		
_	YN		perature:		
Fever	Rash		p Rate:	` ,	·
Respiratory	Sensory (e.g.anosmia, ageusia)	□□ Oxy	gen saturation (%	%) :	in air 🔲 in O ₂ 🔲
Cardiac	☐☐ Headache	□□ Cen	tral cap refill (s):		
Gastrointestinal	Other Neuro			_	
Musculoskeletal	□□ Shock	□□ III aı	opearance:	Y□ N	
Lymphadenopathy	Conjunctivitis				
Oedema/Ascites	☐☐ Mucositis BCG	∐∐ AVF	PU 🔲, GCS 🔲, E	Blantyre ∐:	
Involvement of extremities	reactivation	□□ Abb	rev. Mental Test S	Score	
	ES (tick all that apply)	<u>'</u>		Don't Know	or Not Stated?
	entia; specify:] Allergic	disease; specify:	
Endoc	rine; specify:] Genetic	e, eg T21; specify:	
Sm	oking; n/day:] Recent	surgery; specify:	
Maligna	ancy; specify:]	Hypertension:	
Pulmo	nary; specify:] Organ tr	ransplant; specify:	_
	stinal; specify:	_]	•	
	gical; specify:	_	_	anguinity; specify:	
	rdiac; specify:	<u> </u>	-	Pregnant; Gestation:	
	I; e.g HIV, specify: e.g. HbSS, specify:	<u> </u>	-	ty Scale (Dalhousie):	
				ematurity; Gestation:	
important mstory	not captured above – į	orease use IC	Div codes:		
1.6 SELF-REPORT	ED ETHNICITY (TICK A	S APPLICABLE)			
White European	Black		sian	Middle-Eastern	South American
☐ North/Mid/East	☐ North African (Morocco,Algeria,Tunisia,Libya,Egypt)	☐ South Asia (Indian,Pakistani,Bangla		☐ Turkish	□ Native American
European ☐ South Europe	☐ Sub-Saharan	☐ South East		☐ Arabian Peninsula	
(Merditerranean & Portugal)	☐ Black Caribbean		,Thai,Malay,Indonesian,Filipino	☐ Jewish	│ │
☐ Finnish		(Afghan,Iranian) □ East Asian			☐ Mixed

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DIAMONDS - COVIL					Jy nu	mber:		_ _ -	_ _	_ - = _		
2.1 INVESTIGATIO	ONS, SEVER Default	RITY & TRE Units if different	Bas	seline t with 1	1 st	bloo	search ods/on iission		search oods	blood	esearch ds (at 28 onvales	8
	dd/mm/yy	Ī										
	hh:mm		<u> </u>									
Haematology Haemoglobin	ارم	<u> </u>				Ī						
White cells		ļ	<u> </u>									
Platelets					İ							
Neutrophils	10 ⁹ /L					ĺ						
Lymphocytes						İ						
Monocytes Egginophilo												
Eosinophils Fibrinogen			Ţ									
D-dimer		<u></u>	ģ			<u>j</u>						
	seconds											
Chemistry												
	mmol/L		ļ									
	micromol/L IU/L					ģ						
	nicromol/L											
Albumin												
CRP	mg/L		<u></u>									
PCT	ng/ml					Ĺ						
Ferritin												
Trop-T												
BNP / NT-proBNP LDH						<u></u>						
	IU/L	ļ	ģ			<u>j</u>						
sCD-25												
Blood gas:	ŷ	· · · · · · · · · · · · · · · · · · ·	·			·		·				
Base excess		<u> </u>	<u> </u>			<u> </u>						
	mmol/L kPa		<u> </u>									
Severity	NΓα											
Patient location			ED Ward] I		ED 🗌 Ward 🔲] ICU 🗆		HDU 🗆] IC	טו 🏻
Ventilation		-	Invasive C		NIV 🗌 Ione 🔲			Invasive O2 only	NIV ☐ None ☐] Nor	IV □ ne □
O ₂ Saturation	%											
FiO ₂ / O ₂ Flow	% or L/min			L/	% □ / min□		% □ L / min □		% □ L / min□		L/m	% □ nin□
Heart rate	/min	1	İ		111111-		<u> - / · · · · · · · · · · · · · · · · · ·</u>		<u> </u>			1111
Systolic BP	mmHg											
Respiratory rate	/min											
Capillary refill GCS/AVPU	/15; AVPU	<u>i</u>										
Inotropes ECMO	ļi	<u> </u>	÷	□ No □		.1	□ No □ □ No □	.1] No □] No □		□ No □ □ No □	
ECMO Haemofiltration	<u> </u>			□ No □ □ No □			_ No] No □		□ No □	
Treatment:	Name	e(s):	:		:	1	eived treatment	:		*		
Antibiotic		<u> </u>	Yes [□ No □	J	Yes 🗆	□ No □	Yes 🗌] No 🔲	Yes [□ No □	
Antiviral	·		.i	□ No □			□ No □] No 🔲		□ No □	
Antimalarial				□ No □		i	□ No □	.i] No □	. i	□ No □	
Steroid > 1mg/kg		ļ	<u> </u>	□ No □ □ No □			□ No □ □ No □] No □] No □		□ No □ □ No □	
Immunoglobulin Monoclonal Ab			.ž			.2	_ No	4] No □		□ No □	
Study Treatment	Study			□ No □		. 	□ No □	. 	No 🗆		□ No □	
Other				□ No □		Yes □] No □	Yes 🗆] No 🗆		□ No □	
Maximum CRP (mg/			Max neι	ıtroph	il cou	ınt(10º/L):		Min lymph	ocyte cou	nt(10 ⁹ /L):	•	
3.1 RADIOLOGY R			Yes N	No						Yes N	0	
Chest X-ray:	Had CXR?					X-r	ray: Other:					
	Normal						interstitial	disease]	
	Infiltrates						: Other					
Pneumonia with Ple	n consolidation eural effusions					MR						
Pulr	monary emboli	<u> </u>				Oth	ier					

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3.2 CARDIAC INVESTI	CATIO	ONE		1st ECHO; Date			Worst ECHO; [)ate
Coronary arteries	GAIR	JNO	Normal	Yes No	Unk		Yes No No	Unk 🗍
For largest two CAs			Location		Olik			
Left coronary: LCA; Circumflex:		Long	ez Z-score					
Cx; Left anterior descending:		Internal dia						
LAD; Right Coronary: RCA		mternai dia	meter mm					
Overall Function		М	yocarditis	Yes No No	Unk [Yes No	Unk 🗌
Lo	ow Car	diac Output		Yes No No	Unk [Yes 🔲 No 🔲	Unk 🔲
			ericarditis	Yes No No	Unk [╛	Yes No No	Unk 🗌
		Valvular reg		Yes ∐ No ∐	Unk L		Yes No	Unk 📙
		actional Shor end-diastoli						
	LV	CHG-Glaston	ECG	Normal	1	Block	Normal□	Block□
			200	Arrhythmia [j	Other	☐ Arrhythmia☐	Other□
4.1 VIROLOGY & BACT			muad			D	esults	
Test	Da	rte(s) (ddm		PCR Ct value	///I			All negative □
SARS-CoV-2				est 1:	Test2:	, ig — ig.	Subsequent Test(s)	
			C	Central line Peri	heral]		All negative □
Blood culture					_	•		3,4 4 2
Blood PCR, rapid antigen			F	RAg □ PCR □				All negative □
(bacteriology)	$ \sqcup $			3 — - —				- Jan
			F	PCR 🗌				Negative □
Blood PCR (virology)				_				- 5
			V	Vhite cells N	eutrophil	s %/abs	./ Lymphocytes %/a	abs/
CSF metrics							se/ CSF lactat	
			C	Culture 🗌 RAg/IF	1 PCR [7		All negative □
CSF bacterial						-		<u>-</u>
			F	CR 🗆				Negative □
CSF virology				о <u> </u>				. rogaaro 🗀
				Dipstix (nitrites, leul	ocytes r	rotein)	Microscopy (epithe	lial & white cells)
Urinalysis				Dipotix (minico, icai	tooytoo, p	notonn)	иногозоору (орино	iai a write cons
			(Culture 🗍 RAa 🗆	PCR []		All negative
Urine organism				Jakaro 🗀 Tong 🗅	o L	_		7 til flogative 🖂
			(Culture RAg/IF	l PCR [1		All negative
Nose, throat swab				Januaro 🗀 Turigini 🗅	. O. C	_		/ III riogativo 🖂
			C	Culture ☐ RAg/IF ☐	1 PCR Γ	7		All negative □
Nasopharyngeal aspirate						-		<u>-</u>
			C	Culture RAg/IF	1 PCR Γ	7		All negative □
Stool bacteriology						-		/ga
			F	RAg/IF ☐ PCR ☐				All negative □
Stool virology				ung 🗀 i oit 🗀				7 til Hogativo 🖂
								Negative □
BAL								rtogative 🖂
	-							Negative 🗌
ETT Aspirate								rtogaave 🗀
								Negative □
Serology results								rtogaave 🗀
Other test								
	<u>: </u>		<u>!</u>					
E 4 E 2 CALICATIVE		A NUCRA/C) I	N TIUC II	INFOC Sales	4l-	-4l		
5.1 – 5.3 CAUSATIVE								
Identify pathogens from re	suits c	k cimicai imai	ngs (eg po	silive swab resuil,	or pain	ognomoni	c varicella rash)	
		yes no				yes no		yes no
Evidence of viral aetic	ology		Eviden	ce of bacterial ca	ause		Non-bacterial/vi	iral? 🔲 🔲
Specify virus(es)			Spec	cify bacteria			Specify pathogen.	
opeony viius(es)			Spec	ony baolena			opeony patriogeri.	
\P		· · · · · · · · · · · · · · · · · · ·	D				D (1	····
Virus consistent with some	e but			consistent with so	me but		Pathogen consistent	
NOT all symptoms?	-11			ymptoms?	£ !!		some but NOT all syn	
Virus likely to account for a features of the illness?	all			likely to account of the illness?	for all		Pathogen likely accou all features of illness?	
realures or the limess?			reatures (or trie lilitiess?			an reacures or inness?	

Study number:

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DIAMONDS - COVID		Stu	dy number:	_ -	_ _ -E
Discharged In Admitted to Transferre Date/time of Date/time of Date/time of If died, was Section 1997. Ward of NIV on was No Ceilling of Country of Countr	f death, if died SARS-CoV-2: likely caus (If admitted) 6.3 SEVERIT only	ry gen	Date/Time left ED: Date: D D M N Date: D D M N Date: D D M N Contributed to d (Note: Days = number of mi yes no Days Days Days Days Hae	eath	CU from ED Y N Died in ED Y N Died in ED Y N
DNACPR? Y N					
7.1 SYNDROME CLAS Cardiovascular	Arrhythmia Congestive Heart Failure	ynd 	rome (tick more than one if Cardiac Arrest Central Artery Vasculitis	req	Cardiac Ischaemia Endocarditis
CIT	Myocarditis		Pericarditis Mad/acurra vamiting		Peripheral Artery Vasculitis
GIT Haematological	Mod/severe diarrhoea Deep Vein Thrombosis		Mod/severe vomiting Pulmonary Embolism		Sickle Crisis
Inflammatory	SLE		Kawasaki Disease		Macrophage Activation Synd.
······ai······ai·····	Juvenile Idiopathic Arthritis	H	COVID-related inflammation	\Box	Steven-Johnson Synd.
Lower respiratory	Asthma Exacerbation		Bronchiolitis		Empyema 🔲
tract	Pleural Effusion		Pneumonia		Pneumothorax
	Undefined LRTI		Viral Induced Wheeze		
Musculoskeletal	Arthralgia		Discitis		Myositis 🔲
	Osteomyelitis		Reactive arthritis		Septic arthritis
	Transient synovitis		— ———————————————————————————————————		Marianita
Neurological	CNS Abscess Seizure	님	Encephalitis Stroke / CVA		Meningitis
Pathogen syndromes	COVID19		EBV/Glandular fever		Flu-like illness
r amogon dynaromod	Hand-foot-mouth	H	Hepatitis A	\exists	HSV stomatitis
	Lyme disease		Malaria		Measles ☐
	Mumps		Pertussis		Roseola/HHV6
	Scarlet fever		Staph scalded skin		Varicella 🔲
Sepsis syndromes	Bacteraemia		Febrile neutropaenia		Purpura fulminans
	Septic shock		Severe sepsis	Щ	Toxic shock syndrome
Soft tissue	Abscess		Cellulitis (skin)		Impetigo
	Infected eczema Peri/orbital cellulitis	님	Lymphadenitis Wound infection		Necrotising fasciitis
Surgical	Appendicitis		Bowel obstruction		Mesenteric adenitis
Surgical	Pancreatitis	片	Peritonitis	님	Wesemene adentis
Undifferentiated fever			Fever without source		Fever unknown origin
The state of the s	Post procedural fever				
Urinary tract	Haemolytic uremic syndrome		Pyelonephritis	П	UTI 🖂
URTI/Ear nose throat	Abscess		Croup		Mastoiditis
	Otitis media		Pharyngitis		Sinusitis
	Stomatitis/lip infection		Tonsillitis		URTI (non-specific)
Other infections	CLABSI (line infection)		Conjunctivitis		
Details of above sy	ndromes (eg location, sub-type)		Other synd	Irom	e, give ICD10 code:

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DIAMONDS - COVID	Study number:		-			- 1	ΕĪ	
		 	 	 	 		- 1	

0.4 DDIMARY DUENOTYDE 4:01	ONE	hay (aaa diagram balay) mriari	itialiaa	the besteriel/vival (selected) h								
8.1 PRIMARY PHENOTYPE- tick	KONE	box (see diagram below), priori	itising	the bacterial/viral (coloured) i	oxes							
	yes		yes		yes							
Definite bacterial infection		Viral syndrome - high or no inflammatory markers		Other infection including parasites								
Probable bacterial infection		Probable viral infection		Uncertain – infection or inflammatory syndrome								
Bacterial syndrome - low or no inflammatory markers		Definite viral infection		Inflammatory syndrome – confident diagnosis								
Unknown bacterial or viral infection		Trivial illness: no investigations, treatment, or follow-up		Other cause of illness – specify:								
8.2 SECONDARY PHENOTYPE - responsible for some or all feat		•			so							
	yes		yes									
Other Infection	<u>-</u>											
Initials concurring with final phenotype: Initials not concurring with final phenotype:												

Lopez Z-scores for coronary arteries: http://www.parameterz.com/sites/coronary-arteries

To view the phenotyping algorithm, please use this link: https://imperialcollegelondon.box.com/s/uemqmuzv7vms3ws7c856797ik4cnhhg1

Click here for the Dalhousie Clinical Frailty Scale

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SAMPLE HANDLING RECORD

STUDY	ID DE	TAILS (DI	S-Centre ID-su	bject ID-episod	le; e.g. DI	S-1101-1001-	-E01)	— DIAM V	ND§		
DIS											
Sample ty	уре	Collected		ne of sample coll		Time to	Sample aliq		Storage location		
		Yes No	Time Point (1st,2nd,Dis)	Date	Time	processing (initial storage)		volume and number (if applicable)			
Example: B EDTA	Blood		1st	22-01-2012	23:45	6h (in fridge)	Plasma (100µl): 5 Plasma (250µl): 1 EDTA aliquot (Volume): Cell Pellet (all) (µl): 4 Smart tube (200µl):	Imp Coll, -80°C freezer 9, box 6			
Blood ED	DTA		Tick if same	draw as blood cult	DI (100 l)						
							Plasma (100μl): Plasma (200μl): EDTA aliquot (Volum Cell Pellet (all) (μl): Plasma (100μl):	e):			
DNA							Plasma (250µl): EDTA aliquot (Volum Cell Pellet (all) (µl):				
RNA: PAXger							Whole tube	ne-made' 🔲 ne-made' 🔲			
TEMPU	ıs						Whole tube 🗌 'Hon	ne-made' 🗌			
Serum	1						Serum (100µI): Serum (250µI): Serum (100µI): Serum (250µI): Serum (100µI): Serum (250µI):				
Throat sv											
Stool/red swab											
BAL sam Other:	iple										
Other:											
Sample	hand	lling devia	ntions		<u>-</u>		-	<u> </u>			
-				other deviation)				TP2	DIS		
BLOOD EDTA		•	•	ediately, was it sto to SOP and store			Y N N	Y	Y		
BLOOD SERUM			<u> </u>	g to SOP and store			Y N N	Y N N	Y N		
SMART tube	If the s	sample was i stored at -80	not stored at -80°C	to SOP and store within 15 minutes,	was it kept a	at 4°C for 1 hour	before Y N	n/a n/a	n/a n/a		
PAX tube	Was t	he sample st	ored for a minimur	n of 24 hours at -20	0°C, before s	torage at -80°C?	? Y N N	Y N	Y \ \ \		
Throat swab		•	ored at -80°C with				Y N	Y N	Y N		
Commen	nts/det	ails <i>Please</i>	give details sta	ating clearly wh	ich sample	e type and wha	at time point was a	ffected.			

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