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| --- |
| Inclusion criteria |
| |  |  |  |  | | --- | --- | --- | --- | |  | **Y** |  | **N** | |  |  |  |  | | Patient is aged 18 years or over |  |  |  | |  |  |  |  | |  |  |  |  | | Patient requires testing for SARS-CoV-2 in the opinion of treating clinician |  |  |  | |  |  |  |  | | Patient has presented with acute symptoms of COVID-19 or chest x-ray changes |  |  |  | | or they are asymptomatic, but require testing for other reasons |  |  |  | |  |  |  |  | |
| Exclusion criteria |
| |  |  |  |  | | --- | --- | --- | --- | |  | **Y** |  | **N** | |  |  |  |  | | It is impossible or unsafe to obtain the required research samples |  |  |  | |  |  |  |  | | The patient is a prisoner |  |  |  | |  |  |  |  | | Serial sampling is not feasible for the patient |  |  |  | |  |  |  |  | |
| Consent |
| |  |  |  |  |  |  | | --- | --- | --- | --- | --- | --- | | Type of consent obtained: | | | | | | |  | Tacit/verbal consent |  | Participant written consent |  | Personal consultee\* | |  |  |  |  |  |  | |  | Professional consultee |  |  |  |  | | \*Relationship to participant | | | | | | |  | | | | | |  |  |  |  |  |  |  |  |  |  |  |  |  | | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | | Date consent obtained: | |  |  | / |  |  | / |  |  |  |  | |  | | D | D |  | M | M |  | Y | Y | Y | Y | | Consent obtained by: |  | | | | | | | | | | | | |
| CRF completion |
| |  |  | | --- | --- | | CRF completed by: |  |  |  |  |  |  |  |  |  |  |  |  |  | | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | | Date completed: |  |  | / |  |  | / |  |  |  |  | |  | D | D |  | M | M |  | Y | Y | Y | Y | |