|  |
| --- |
| 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 |

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