

INSERT  
MRN and NHS Number

# Patient Consent Form

## Barts BioResource Health Data and Sample(s) donation

(such as blood/saliva/swabs/urine/faeces and/or soft/solid tissue samples)

**REC reference:** 14 / EE / 0007

**REC reference:** 21 / EE / 0284

### Patient Consent Form

#### Donation of Health Data and Sample(s) to the Barts BioResource.

The study has been explained to me by:

Prof/Dr/Mr/Mrs/Ms \_\_\_\_\_

**Participants should complete the whole of this form themselves.**

| Please initial each box as appropriate   | YES | NO |
|--|-----|----|
| <p>I confirm that I have read and understand the <b>Barts BioResource</b> Patient Information Sheet (Barts BioResource Health Data and Sample(s) donation (such as blood/saliva/swabs/urine/faeces and/or tissue sample), v11.0; Dated 1<sup>st</sup> January 2022 for the above study. I understand that my medical information will be treated confidentially.</p> <p>I agree to my healthcare related data (appropriately de-identified) being included on publicly accessible Web sites to facilitate research, teaching and education of healthcare and disease. I understand that relevant sections of my health data collected during the study may be looked at by individuals from regulatory authorities, academic research partners, the NHS and associated delivery organisations, where it is relevant to my taking part in this research.</p> <p>I give permission for these individuals to have access to my records. I understand that I can ask for more information at any time using the contact details on the Information Sheet.</p> <p>I understand that information held by the NHS (e.g. hospitals, NHS Digital, Healthcare Quality Improvement Partnership (HQIP)), my General Practitioner (GP) and records maintained by the Office for National Statistics (ONS) may be used to follow up my health status, I give permission for this information to be obtained by the research team if necessary.</p> |     |    |
| I agree to be contacted in the future for further information on clinical studies and trials for healthcare research.  |     |    |



Study Role: \_\_\_\_\_

**If the subject was assisted during the Consent process please complete:**

The consent form was read to the subject, and the person signing below acted as an interpreter/ translator for subject during the consent process. The person signing below attests that the study was accurately explained and understood by the subject.

Please provide the language used to take consent: \_\_\_\_\_

Signature of Person Assisting in the Consent Discussion

\_\_\_\_\_  
Signature

\_\_\_\_\_  
Printed name

\_\_\_\_\_  
Date (DD/MM/YYYY)