





Please <u>initial</u>

box. Do

## **CONSENT FORM**

Plasma biomarkers in stratifying patients referred via the lower gastro-intestinal (LGI) suspected cancer two-week wait (2WW) pathway

MOTION study

IRAS: 321809

Participant ID:

|   | <u>not tick</u> |
|---|-----------------|
| Please read the statements below. Put <u>initials</u> of your name in the boxes if you agree, put an X in any box where you do not agree and sign the form. <b>Use BLACK pen only</b> |                 |
|   |                 |
| 1. I confirm that I have read and understand the Patient Information Sheet dated [DATE] (version  |                 |
| <b>[VERSION NUMBER]</b> ) for MOTION study. I have had the opportunity to consider the information, ask   |                 |
| questions and have had these answered satisfactorily.   |                 |
| 2. I understand that my participation is voluntary and that I am free to withdraw at any time, without  |                 |
| giving any reason and without my care or legal rights being affected.   |                 |
| 3. I agree that confidential study data collected prior to my withdrawal from the MOTION study can be   |                 |
| used in the study reports. I understand that if I lose capacity to consent during the study I will be   |                 |
| withdrawn and the agreed data already collected can be used.  |                 |
| 4. I understand that relevant sections of my medical notes and data collected during the MOTION study   |                 |
| may be looked at by individuals from the Queen Mary, University of London (QMUL) or Barts Health  |                 |
| NHS Trust (BHT). I give permission for these individuals to have access to my records.  |                 |
| 5. I agree for my contact details and identifiable personal data to remain within the study site (QMUL or   |                 |
| BHT). I understand that coded personal data, including data from medical records and study-specific   |                 |
| data, may be shared with collaborators within the EU for the purposes of research and data analysis.  |                 |
| 6. I agree to my GP or/ and referring doctors being informed of my participation in the MOTION study.   |                 |
| 7. I understand that the study data collected will be used for medical research only. I will be given a   |                 |
| Unique Identification Number (UIN/ Patient ID) in order to ensure that my data is confidential and will not   |                 |
| directly identify me.   |                 |
| 8. I agree my confidential study data can be used in the analysis and reporting of the study findings.  |                 |
| 9. I understand that participants will be asked to give a 20 millilitres of blood sample, to be taken in clinic   |                 |
| or in endoscopy or in the radiology department. I consent to having my blood drawn.   |                 |
| 10. I understand that a laboratory will analyse my blood sample and the MOTION study team will record   |                 |
| this information.   |                 |
| 11. I understand that a part of my blood sample will be transported to our collaborators in France.   |                 |
| A portion of the blood sample collected for this study will be sent to Progastrin Manufacturing, Biodena  |                 |
| Care, located at Cap Sigma-Zac Euromedicine II, 1682 Rue de la Vasière, 34790 Grabels, France, for  |                 |
| the purpose of biomarker analysis.  |                 |
| 12. I understand that this MOTION study also involves the analysis of my DNA and RNA from plasma  |                 |
| and I agree for genetic analysis to be done.  |                 |
| 13. I understand that the storage of my samples will be pseudonymised (labelled with an ID number   |                 |

instead of personally identifying information) and outcomes of the analysis are unlikely







| to have any implications for me personally.  |                  |                                |       |
|--|------------------|--------------------------------|-------|
| 14. I understand that my contact details will be kept in order for the study team to communicate with me |                  |                                |       |
| about the study and for QMUL to retain study-related information for 5 years after the study ends, after |                  |                                |       |
| which point all study materials will be destroyed.   |                  |                                |       |
| 15. I understand that relevant sections of my medical notes and data collected during                    |                  |                                |       |
| the study may be looked at by individuals from QMUL or BHT, from regulatory authorities or               |                  |                                |       |
| from the NHS Trust, where it is relevant to my taking part in this research. I give permission for       |                  |                                |       |
| these individuals to have access to my records.  |                  |                                |       |
| 16. I agree to take part in the study.   |                  |                                |       |
| 17. OPTIONAL   |                  |                                |       |
| I agree to be contacted by the study team to complete a survey at the end of the two-week wait           |                  |                                |       |
| pathway using my contact details.  |                  |                                |       |
| 18. OPTIONAL   |                  |                                |       |
| I prefer to be informed and receive information about this questionnaire through the following           |                  |                                |       |
| methods.   | □ phone intervie | w □ SMS                        |       |
|  |                  |                                |       |
| 19. OPTIONAL   |                  |                                |       |
| I understand that the information collected about me will be used to support other research in the       |                  |                                |       |
| future and may be shared anonymously with other researchers.   |                  |                                |       |
| 20. OPTIONAL   |                  |                                |       |
| Following the present study, any residual (left-over) plasma may be stored and used by the research      |                  |                                |       |
| team at QMUL, for future research studies. Any future studies will have Ethics Committee permission.     |                  |                                |       |
| 21. OPTIONAL   |                  |                                |       |
| I agree to be contacted about future related stud  | ies.             |                                |       |
| Print Name of Participant:   | Date:            | Participant's Signature:       |       |
|  |                  |                                |       |
|  |                  |                                |       |
| Print Name of person taking consent:   | Date:            | Signature of person taking con | sent: |
|  |                  |                                |       |
|  |                  |                                |       |