



Hepatitis B in the Mongolian Community in London: An investigation of epidemiology and burden of disease to inform clinical and public health interventions (Hep-MOLO)

Name of Chief Investigator: Prof Philippa Matthews

IRAS Project ID: 341947

Participant Study ID: M O L O - - - - -

Initial box

Part 1	
I confirm that I have read and understand the participant information sheet for the above study and have had the opportunity to ask questions which have been answered fully.	
I give my consent to take part in clinical screening for blood borne infections, metabolic risk factors and liver disease.	
I am happy to take a questionnaire about my liver health. I understand I do not have to answer all the questions, and the data will be stored anonymously and analysed for research purposes.	
I give my consent to be contacted with my results either by telephone, post or email.	
I give my consent for my general practitioner to be contacted with my results.	
I understand that if I am diagnosed with chronic viral hepatitis, clinical data e.g. liver test results, appointment attendance, may be collected up to 1 year following diagnosis for this study. Clinical records will only be accessed by authorised members of the clinical team involved in the research study and anonymised before analysis.	
I understand that anonymised data collected in the study may be looked at by authorised individuals from the clinic team, the Sponsor (UCL), NHS bodies, by people working on behalf of the Sponsor, or by representatives of regulatory authorities where it is relevant to taking part in this research. I give permission for these individuals to have access to these records.	
I give permission for the research team to collect, store, analyse and publish anonymous research data obtained from my participation in this study. I understand that my personal details will be kept confidential.	
I understand that my participation is voluntary and that I am free to withdraw at any time, without giving any reason, and without my medical care or legal rights being affected. I understand that should I withdraw, the information collected so far cannot be deleted and that this information may still be used in the study analysis.	
Part 2 (optional)	
I give my consent for additional blood bottles to be collected and stored as part of research into chronic viral hepatitis and metabolic risk factors (for example, analysis of blood fats, immune system function and viral factors).	
I understand that my blood samples may be tested for genetic factors which might influence my immune system.	
I understand that these blood tests are for research purposes only and I will not be informed of my individual results.	
I understand that my blood samples will be stored at the Francis Crick Institute, or collaborating laboratories, for up to 10 years, and may be used in further studies following appropriate ethical approval.	
Part 3 (optional)	
I consent to being contacted for further research within the scope of this project (e.g. interview or focus group). I understand I am under no obligation to take part.	
Once the study has ended, I would like to know what the study found out and agree for my personal details to be used for sending me this information.	

Name of Participant

Signature

Date

Name of Person taking consent

Signature

Date