

Informed Consent Form

Assessing gut permeability using spectroscopy

Study title: Non-invasive transcutaneous spectroscopy for the assessment of gut permeability (GutPerm)

IRAS Project ID: 242462

Patient Trial ID Number: _____

Name of Principal Investigator: Dr Alex Thompson

Name of patient to whom this ICF applies: _____

Please initial boxes:

1. I confirm that I have read and understand the Patient Information Sheet "Assessing gut permeability using spectroscopy v4.0 31-May-2018" for the above study and have had the opportunity to ask questions about the study and understand what is involved.

2. I understand that my participation is voluntary and that I am free to withdraw at any time, without giving any reason, without my medical care or other legal rights being affected, and I understand that data collected up to the point of my withdrawal may still be used.

3. I agree to have a 'Spectroscopic gut permeability test' and understand that there is a small risk of nausea and other allergic reactions (including anaphylaxis) with this test.

4. I understand that sections of any of my medical notes may be looked at by authorised individuals from Imperial College Healthcare NHS Trust, from Imperial College London, or from regulatory authorities where it is relevant to my taking part in this research. I give permission for these individuals to have access to my records.

5. I agree to allow information about me to be collected, analysed, reported and transferred to other approved collaborators within and outside the European Economic Area for healthcare and/or medical research purposes. I understand that my identity will remain anonymous.

6. I wish to be contacted by e-mail with a summary of the research findings once the study is complete. Your e-mail address will not be used for any other purpose.

Contact e-mail: _____

If you do NOT wish to be contacted regarding the study results, please put a line through all of point 6 above, do NOT leave a contact e-mail address and do NOT initial the box as indicated – you can still participate in the main trial.

7. I agree to take part in the study.

Please sign and date overleaf.

Name of Subject

Signature

Date

Name of Person taking consent

Signature

Date

1 copy for subject; 1 copy for Principal Investigator; 1 copy to be kept with hospital notes