

Blood Biomarkers for diagnosis and prognosis of concussion

CONSENT FORM

NCT NUMBER: NCTXXXXXXXXX

08/MAY/2019

Study Title: Blood Biomarkers for diagnosis and prognosis of concussion

INFORMED CONSENT FORM

Subject Number:

Principal Investigator: Dr John Mulvihill

Participation in this study is voluntary and you may withdraw at any time for any reason

	<i>Please tick appropriate box</i>	
The research project and procedure associated with it have been fully explained to me.	YES <input type="checkbox"/>	NO <input type="checkbox"/>
All experimental procedures have been identified and no guarantees have been given about the possible results.	YES <input type="checkbox"/>	NO <input type="checkbox"/>
I have had the opportunity to ask questions concerning any and all aspects of the project and any procedures involved.	YES <input type="checkbox"/>	NO <input type="checkbox"/>
I am aware that participation is voluntary and I may withdraw consent at any time.	YES <input type="checkbox"/>	NO <input type="checkbox"/>
I am aware that my decision not to participate or to withdraw will not restrict my access to health care services normally available to me.	YES <input type="checkbox"/>	NO <input type="checkbox"/>
Confidentiality of records concerning my involvement in this project will be maintained in an appropriate manner and my data is protected.	YES <input type="checkbox"/>	NO <input type="checkbox"/>
I understand that portions of my team medical records may be looked at by responsible individuals from the research team. I give permission for these individuals to have access to my records.	YES <input type="checkbox"/>	NO <input type="checkbox"/>
I understand that the investigators have such insurance as is required by law in the event of injury resulting from this research	YES <input type="checkbox"/>	NO <input type="checkbox"/>

Analyses of all samples and information collected will be conducted in the University of Limerick. On some occasions the analyses may be done in collaboration with third parties, including commercial partners, which may require samples to be shipped to and/or analysed by these organisations. In all cases, samples and data will be coded with anonymized identifier numbers.

(please tick box if you agree) ☐

I am happy that the samples and data will be stored and used in other research studies within this topic of research (i.e. neurological disorders).

(please tick box if you agree) ☐

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Access to my samples and/or my data will require approval from the Executive Management Group of the Faculty of Science and Engineering, University of Limerick.

(please tick box if you agree) ☐

Any identified biomarkers from this study can be used and disseminated by the principal investigator.

(please tick box if you agree) ☐

I agree that my contact details may be made available for a follow up check on my status following my participation in this project or its Clinical Research Ethics Committee approved collaborators.

(please tick box if you agree) ☐

I, the undersigned, hereby consent to participate as a subject in the described project conducted at the University of Limerick. I have received a copy of this consent form and a copy of the patient information leaflet for my records. I understand that if I have any questions concerning this research, I can contact **Dr. John Mulvihill, Biomedical Engineer, University of Limerick.**

Telephone number: (061) 237719.

Subject's Signature: _____

Date: _____
dd mon yy

NAME (BLOCK LETTERS): _____

Time: _____

Researcher's Signature: _____

Date: _____
dd mon yy

NAME (BLOCK LETTERS) _____

Time: _____