RESEARCH PARTICIPANT INFORMED CONSENT FORM

GENERAL DATA PROTECTION REGULATION ADDENDUM

As required by the law of the European Union and its member countries, and/or the related law of the United Kingdom, additional information is being provided to you as a study participant.

For purposes of EU and UK law, particularly the General Data Protection Regulation (GDPR) and UK GDPR, the lawful basis for the use of your data is your consent via this form, as well as the public interest in the research being conducted.

The Consent Form indicates whether the research team will collect Personal Information about you. In addition to any basic information that may identify you, Personal Information may also include the results of any tests, surveys or procedures described in the informed consent form.

Any such Personal Information will be treated in compliance with applicable data protection laws. In addition to the uses of your information shown on the Consent Form, your Personal Information may be used to:

- protect your vital interests (for example, for purposes of monitoring an epidemic or other public health emergency); and
- answer your data protection requests (if any).

In addition to entities listed in the Consent Form, your Personal Information may be shared with entities or organizations in a country (including the United States) that have not received an adequacy decision by the European Commission, based on your consent via this form, and the necessity of the transfer for the public interest. An “adequacy decision” is a determination by the European Union that a particular non-EU country’s laws or other international commitments ensures an adequate level of protection of personal data.

You may have additional rights with respect to your Personal Information according to the EU or UK GDPR. If you wish to exercise any of the rights described below, please contact a member of the study team identified in the Consent Form, or the Privacy Office.

- You have the right to see the information being collected about you in the study. To ensure integrity of the study, you may not be able to review some of the data until after the study has been completed.
- You have the right to correct or update your Personal Information if it is inaccurate.
- You have the right to limit the collection and use of your Personal Information under certain circumstances (for example, if you think that the information is inaccurate).

- You have the right to withdraw from the study. If you withdraw from the study, you will no longer be able to participate in the study. No new information or samples will be collected about you or from you by the study team. Your withdrawal has no effect on the lawfulness of the data processing that occurred prior to your withdrawal.
- You have the right to receive your Personal Information in a structured, common computer format (for example, in a readable text electronic file or chart) for your own purposes or for giving it to others, as required by applicable data protection laws. You may not have the right to receive your Personal Information that has been used for public interest purposes (for example, for reporting incidents of disease to public health officials) or in the exercise of official authority (for example, responding to information requests from public agencies or monitoring drug safety)
- You have the right to request the deletion of your Personal Information if you are no longer participating in the study. However, there are limits on your ability to request deletion of your Personal Information, for example, after the data has been de-identified, and any identifiers or links have been destroyed.
- You have the right to file a complaint with a data protection authority (http://ec.europa.eu/justice/data-protection/article-29/structure/data-protection-authorities/index_en.htm).
- You may contact the UF Data Protection Officer with any questions about the study. Please call the UF Privacy Office to reach the GDPR Data Protection Officer.