## Temporary Addendum to Informed Consent

## Title of research study: ***[insert title of research study here. This should be identical to the protocol title used in Cayuse IRB]***

## IRB #: ***[IRB-FY20XX-XX]***

## Investigator: ***[insert name of principal investigator]***

## Overview:

## In light of the current global pandemic and the risks to human health and safety stemming from direct personal contact, the Villanova Institutional Review Board (IRB), which provides ethical review and oversight for human subject research, has mandated that all human subjects research be conducted remotely rather than in-person. The following section describes the changes you will experience from what was originally expected and described in the main consent document as a result of that IRB requirement.

* Instead of [describe first activity/experience and how it will change]
* Instead of [describe second activity/experience and how it will change]
* Use as many bullet points as necessary to clearly explain all deviations from the main consent document. You may adapt this format as necessary to describe your changes, as long as the participant will be able to fully understand the expected differences.

## Participant Acknowledgement

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| Your signature here documents that you have read the changes described above and understand that they outline the experience you will have during the temporary remote research period for this project. |
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| Signature of subject |  | Date |
|  |  |

Printed name of subject