**DIRECTIONS**

 **Consent Form**

*This document is an example. There are various ways to set up this document. This is just one example for you to use.*

* Read the consent form and adapt according to your research.
	+ Adapt **ORANGE TEXT** to fit your research.
	+ Choose the appropriate statement in **GREEN TEXT** that applies to your research. Delete the other statement not being used.
	+ Choose the appropriate **BLUE TEXT** option. Delete the other option not being used.
	+ Deletethe **RED TEXT** from the consent form.
* Format the document to un-bold the text and change the font color to black.
* Expedited and Full Board IRBs **REQUIRE** a consent form, signature, and date. Unless, a waiver of written consent is requested through the IRB application. When the IRB grants the waiver of written consent the researcher will remove the signature and date line from the consent form.
* If your research is exempt, then refer to the information sheet. Exempt research does **NOT** require a consent form or a signature line.

**CONSENT FORM**

**{Title of your study}**

**What is this research studying?**

This study will help us learn how **{briefly describe the purpose of your study.}**

**What would I do if I participate?**

In this study, you will be asked to **{describe what their participation entails in this study.} {If you are providing an incentive/compensation to the participant provide the following: a description of the incentive/compensation; any restrictions or requirements for participant compensation, if it is a drawing the odds of winning, and the time frame of when the participant will receive the incentive/compensation.}**

**Can I quit if I become uncomfortable?**

Yes, absolutely. **{TTU PI’s Name}** and Texas Tech University’s Institutional Review Board have reviewed this research project and think you can participate comfortably. However, you can skip parts of the research you are not comfortable with and stop at any time. You will keep all the benefits of participating even if you stop. Participating is your choice.

**How long will participation take?**

We are asking for **{amount of time}** of your time.

**How are you protecting privacy?**

Your name will not be linked to any material in reports, publications or presentations. No one other than the researchers associated with this project will have access to the raw data. All related documentation will be stored in the researcher’s locked office **{and/or}** on a password protected computer.

**What will happen to my data?**

**Identifiers might be removed from the {CHOOSE ONE OPTION: identifiable private information OR identifiable biospecimens} and after such removal the {CHOOSE ONE OPTION: information or biospecimens} could be used for future research studies or distributed to another investigator for future research studies without additional informed consent from you or your legally authorized representative.**

**or**

**Your {CHOOSE ONE OPTION: information OR biospecimens} collected as part of the research, even if identifiers are removed, will not be used or distributed for future research studies.**

**What are the benefits and risks of participating in this research?**

**{CHOOSE ONE OPTION: There are no anticipated risks or benefits to your participation in this research. OR The risks associated with this research are (Enter risk(s)). and The benefits associated with this research are (Enter benefit(s)).}** We appreciate your time and effort with this research study.

**I have some questions about this study. Who can I ask?**

The study is being run by **{TTU PI’s Name and Co-Investigator’s Name}** from the Department **Department’s Name** at Texas Tech University. If you have questions, you can call **{him/her}** at **{XXX-XXX-XXXX}**.

Texas Tech University also has an Institutional Review Board that protects the rights of people who participate in research. You can contact them at 806-742-2064 or hrpp@ttu.edu.

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Signature of Participant Date

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Printed Name of Participant