**These are examples of a concise summary.**

**The content must be modified to be applicable to your study.**

**This concise summary information appears as the first portion of a consent form with the complete consent form content to follow.**

**The concise summary can appear as part of the consent document itself and does not need to be a standalone document.**

**Delete yellow highlighted text and header/footer.**

**Adjust formatting as needed.**

**Example 1**

**Concise Summary for Adult Consent**

**Important Information You Need to Know:**

* The purpose of this research study is to (*state the purpose*).
* Participation to this research is voluntary. You will complete (*state the research activity that the participant will do. E.g. a computer-based task/an online survey/an in-person individual interview/phone interview/will be observed, etc.*).
* This will be a/an (*state the method. E.g. one-time survey/pre and post survey/audio-recorded interview/video-recorded observation, etc.*).
* This will require (*state the duration. E.g. 15 minutes, one hour, or a total of 3 hours*) of your time (*make sure to give details of each activity if it includes multiple visits*).
* Follow-up (*e.g interview might occur if clarifications are needed/survey 8 weeks after completion of the program will be sent via email*). Total study duration is about (*state the duration e.g. 6 months/1 year*).
* The greatest risks of this study include the possibility of (*state the risk e.g. emotional distress/embarrassment*) during the completion of survey or during the interview with questions involving (*e.g. state the types of questions that will be asked e.g. your mental health/eating habits*) and loss of confidentiality.
* Potential benefits of participation include (*e.g. potential reduction in symptoms, gaining skills, professional development, etc.)*

If you are interested in learning more about this study, please continue to read below.

**Example 2**

**Concise Summary for Adult Consent**

The purpose of this research study is to (*state the purpose*). Participation to this research is voluntary. Participants will complete (*state the research activity that the participants will do. E.g. a computer-based task/an online survey/an in-person individual interview/phone interview/will be observed, etc.*). This will be a/an (*state the method. E.g. One-time survey/pre and post survey/audio-recorded interview/video-recorded observation, etc.*). This will require (*state the duration. E.g. 15 minutes, one hour, or a total of 3 hours*) of your time (*make sure to give details of each activity if it includes multiple visits*). Follow-up (*e.g interview might occur if clarifications are needed/survey 8 weeks after completion of the program will be sent via email*). Total study duration is about (*state the duration e.g. 6 months/1 year*).

The greatest risks of this study include the possibility of (*state the risk e.g. emotional distress/embarrassment)* during the completion of survey or during the interview with questions involving (*e.g. state the types of questions that will be asked e.g. your mental health/eating habits*) and loss of confidentiality. Potential benefits of participation include (*e.g. potential reduction in symptoms, gaining skills, professional development, etc.*)

If you are interested in learning more about this study, please continue to read below.