**INSTRUCTIONS**: **Use this template to structure your consent form. Text in brackets and capitals are instructions for content for each section and should be deleted. The text in each section is provided as a suggestion. Some of the information in this form may not be relevant to your study and can be deleted. You should also add any information that applies to the study but is not in this template.**

**DELETE THIS BOX PRIOR TO SUBMITTING YOUR CONSENT FORM TO THE UHSRC.**

**Parental Consent Form**

Project Title:

Principal Investigator: [Name, institutional affiliation]

Co-Investigator: [IF APPLICABLE][Name, institutional affiliation]

Faculty Advisor: [ONLY FOR STUDENT PI] [Name, institutional affiliation]

Sponsor: [IF APPLICABLE]

**Invitation to participate in research**

Your child is invited to participate in a research study. In order to participate, your child must [LIST ELIGIBILITY CRITERIA]. Participation in research is voluntary. Please ask any questions you have about participation in this study

**Important information about this study**

* The purpose of the study is [BRIEFLY DESCRIBE STUDY PURPOSE].
* Participation in this study involves [BRIEFLY DESCRIBE STUDY PROCEDURES AND TASKS].
* Risks of this study include [BRIEFLY DESCRIBE STUDY RISKS].
* The investigator will protect your confidentiality by [BRIEFLY DESCRIBE DATA STORAGE PROCEDURES].
* Participation in this research is voluntary. Your child does not have to participate, and if you and your child decide to participate, you or your child can stop at any time.

**What is this study about?**

The purpose of the study is [DESCRIBE STUDY PURPOSE]

**What will happen if my child participates in this study?**

Participation in this study involves

* [LIST STUDY PROCEDURES HERE]
* [INDICATE ANY PROCEDURES THAT ARE EXPERIMENTAL]
* [IF THERE ARE MULTIPLE STUDY VISITS, DESCRIBE WHAT HAPPENS AT EACH VISIT USING SEPARATE SUB-SECTION HEADINGS: VISIT 1, VISIT 2, ETC.]
* [EXPLAIN HOW LONG PARTICIPATION LASTS, OVERALL AND BROKEN DOWN BY DAYS/SESSIONS/ETC.]

[IF CLINICALLY RELEVANT RESULTS WILL BE SHARED, MODIFY TEXT IN CAPS AND ADD: We may learn information about your child’s health as a part of this research. We WILL/WILL NOT share this information with you. EXPLAIN HOW INFORMATION WILL BE RETURNED, IF APPLICABLE]

[IF AUDIO/VIDEOTAPING, MODIFY TEXT IN CAPS AND ADD: We would like to AUDIO/VIDEO record your child for this study. If your child is AUDIO/VIDEO recorded, it will be possible to identify him or her through his or her VOICE/IMAGE. If you agree to allow your child to be AUDIO/VIDEO recorded, sign the appropriate line at the bottom of this form.]

[IF RANDOMIZATION WILL OCCUR, MODIFY THE TEXT IN CAPS AND ADD: Your child will be assigned by chance (like the flip of a coin) to one of [XX] groups. One group will [DESCRIBE GROUP PROCEDURES] and the other group will [DESCRIBE GROUP PROCEDURES. ADD DESCRIPTIONS FOR EACH GROUP]. You, your child, or the investigator cannot choose which your group. Your child has an equal chance (1 out of XX) of being assigned to either study group.]

[IF BIOLOGICAL SAMPLES ARE COLLECTED, MODIFY THE TEXT IN CAPS AND ADD: This research WILL/WILL NOT involve whole genome sequencing. Please ask the investigator any questions you have about how your child’s genetic information will be used.]

**What are the anticipated risks for participation?**

[LIST RISKS HERE. YOU MAY CHOOSE FROM SAMPLE TEXT BELOW AND ADD ADDITIONAL TEXT. TIME AND INCONVENIENCE ARE NOT RISKS.]

There are no anticipated physical or psychological risks to participation.

The primary risk of participation in this study is a potential loss of confidentiality.

Some of the [SURVEY/INTERVIEW/FOCUS GROUP] questions are personal in nature and may make your child feel uncomfortable. Your child does not have to answer any questions that make him or her feel uncomfortable or that he or she does not want to answer.

**Are there any benefits to participating?**

[LIST BENEFITS HERE. MOST STUDIES WILL NOT DIRECTLY BENEFIT SUBJECTS. USE SAMPLE TEXT BELOW AND ADD ADDITIONAL TEXT. COMPENSATION OR LEARNING ABOUT ONESELF OR THE RESEARCH PROCESS ARE NOT BENEFITS.]

You and your child will not directly benefit from participating in this research.

Benefits to society include [LIST BENEFITS TO SOCIETY].

**How will my child’s information be kept confidential?**

[EXPLAIN MEASURES YOU WILL USE TO KEEP DATA SECURE. YOU MAY CHOOSE FROM THE SAMPLE TEXT BELOW AND ADD ADDITIONAL TEXT]

We plan to publish the results of this study. We will not publish any information that can identify your child. [IF IDENTIFIABLE DATA WILL BE PUBLISHED: We would like to publish your child’s identifiable information, with your permission. We will ask you to initial a statement at the bottom of this form to give us permission to identify your child in publications. Please note that once your child’s identifiable information is published, it cannot be removed, even if you and your child are unhappy with the publication.]

We will keep your child’s information confidential by [STATE CONFIDENTIALITY MEASURES: USING A CODE TO LABEL DATA WITH THE CODE LINKED TO IDENTIFIABLE INFORMATION IN A KEY STORED SEPARATELY FROM DATA, NOT COLLECTING ANY IDENTIFIABLE INFORMATION (INCLUDING YOUR NAME), ETC.]. Your information will be stored in a [STATE HOW DATA WILL BE STORED: PASSWORD-PROTECTED FILE ON A PASSWORD-PROTECTED COMPUTER, LOCKED FILING CABINET, ETC.].

We will make every effort to keep your child’s information confidential, however, we cannot guarantee confidentiality. Other groups may have access to your child’s research information for quality control or safety purposes. These groups include the University Human Subjects Review Committee, the Office of Research Development, the sponsor of the research, or federal and state agencies that oversee the review of research, including the Office for Human Research Protections and the Food and Drug Administration. The University Human Subjects Review Committee reviews research for the safety and protection of people who participate in research studies.

[IF MANDATORY REPORTER, ADD: If, during your child’s participation in this study, we have reason to believe that elder abuse or child abuse is occurring, or if we have reason to believe that your child is at risk for being suicidal or otherwise harming him or herself or anyone else, we must report this to authorities as required by law. We will make every effort to keep your child’s research information confidential. However, it may be possible that we have to release your child’s research information. If this were to occur, we would not be able to protect your child’s confidentiality.]

[IF FOCUS GROUP OR GROUP INTERVIEW, MODIFY TEXT IN CAPS AND ADD: The investigators will ask your child and the other people in the group to use only first names during the FOCUS GROUP/INTERVIEW session. The investigators will also ask your child not to tell anyone outside of the group about anything that was said during the group session. However, we cannot guarantee that everyone will keep the discussions private.]

**Storing study information for future use**

We WILL/WILL NOT store your child’s information to study in the future. Your child’s information will be labeled with a code and not your child’s name. Your child’s information will be stored in a password-protected or locked file.

We may share your child’s information with other researchers without asking for your permission, but the shared information will never contain information that could identify you or your child.

**What are the alternatives to participation?**

[LIST ALTERNATIVES HERE. YOU MAY USE SAMPLE TEXT BELOW. IF COURSE CREDIT IS OFFERED FOR PARTICIPATION, LIST ALTERNATIVE METHODS OF OBTAINING SUCH CREDIT – COMPLETING ALTERNATE ASSIGNMENT, ETC.]

The alternative is not to participate.

The study treatment offered in the study is experimental in nature and is not available outside of this research study. If you would like to seek alternate, standard of care treatment for your child, the investigator can provide you with resources.

**Are there any costs to participation?**

[LIST COSTS OF PARTICIPATION. TIME IS NOT A COST]

Participation will not cost you or your child anything.

You will be expected to pay for the study treatment provided at a reduced rate of $[XX] per session. You are being charged for study treatment because the treatment used is considered standard of care and is accessible whether or not your child participates in this research study.

**Will my child be paid for participation?**

[DESCRIBE COMPENSATION FOR THE STUDY. YOU MAY USE THE SAMPLE TEXT PROVIDED BELOW OR ADDITIONAL TEXT. REFRESHMENTS DO NOT CONSTITUTE COMPENSATION. INCLUDE INFORMATION ABOUT COURSE CREDIT IF APPLICABLE]

Your child will not be paid to participate in this research study.

Your child will be paid $[XX] for completing this study. If your child does not complete this study, your child will be paid $[XX] for each hour of participation.

Your child will be given a gift card for $[XX] for participating in this research study.

Your child’s school will receive [DESCRIBE INCENTIVES OR COMPENSATION FOR SCHOOL IF APPLICABLE].

**What happens if my child is injured while participating in the research? [THIS SECTION IS OPTIONAL. IF THE STUDY IS NOT GREATER THAN MINIMAL RISK OR IF PHYSICAL INJURY IS NOT LIKELY, REMOVE THIS SECTION]**

If your child is injured as a result of participating in this study, we will assist you and your child in getting necessary medical treatment. You or your insurance company will be responsible for the cost. Eastern Michigan University does not provide any form of compensation for injury.

**Who will profit from the study results? [THIS SECTION SHOULD ONLY BE USED IF BIOLOGICAL SPECIMENS WILL BE COLLECTED AND RETAINED]**

The biological specimens and/or your personal information may be used by the investigator and by Eastern Michigan University for commercial profit. If this happens, you will not be notified. You will not share in the profit.

**Study contact information**

If you or your child has any questions about the research, you can contact the Principal Investigator, [PI NAME], at [EMAIL ADDRESS] or by phone at [PHONE NUMBER]. [STUDENT INVESTIGATORS MUST ADD THE FOLLOWING TEXT: You can also contact [NAME]’s adviser, [ADVISER’S NAME], at [ADVISER EMAIL ADDRESS] or by phone at [ADVISER PHONE NUMBER].

For questions about your child’s rights as a research subject, contact the Eastern Michigan University Human Subjects Review Committee at human.subjects@emich.edu or by phone at 734-487-3090.

**Voluntary participation**

Participation in this research study is your and your child’s choice. Your child either will be asked independently for assent or his or her dissent will be respected. You and your child may refuse to participate at any time, even after signing this form, with no penalty or loss of benefits to which you and your child are otherwise entitled. You and your child may choose to leave the study at any time with no loss of benefits to which you and your child are otherwise entitled. If you and your child leave the study, the information your child provided will be kept confidential. You and your child may request, in writing, that your child’s identifiable information be destroyed. However, we cannot destroy any information that has already been published.

**Statement of Consent**

I have read this form. I have had an opportunity to ask questions and am satisfied with the answers I received. I give my consent to for my child to participate in this research study.

**If you would do not agree to let your child participate in this study, contact the investigator at [PI PHONE] or [PI EMAIL] by [ADD CUT-OFF DATE FOR REFUSAL TO PARTICIPATE]. If you do not contact the investigator by [CUT-OFF DATE FOR REFUSAL TO PARTICIPATE], your consent will be assumed.**