## Informed Consent for Participation in a Research Study

## Title of Research Study: [insert title of research study here]

## Investigator: [insert name of Principal Investigator and Department]

## Key Information:

## {Should not exceed one page}

## You are being asked to take part in a research study about … (include general description of study). This page will give you key information to help you decide whether or not you want to participate in this study. More detailed information can be found on the next pages. Ask the research team questions during the consent process, and use the contact information on this form to ask questions later.

**WHAT IS THE PURPOSE, PROCEDURES, AND DURATION OF THIS STUDY?**

***Briefly*** *describe the purpose if the study and the procedures to be followed in lay terms. Put detailed descriptions in the main consent document (next pages).*

By doing this study, we hope to learn …Your participation in this research will last about *{hours, days, months, years}.*

**WHAT ARE THE REASONS YOU MIGHT CHOOSE TO VOLUNTEER FOR THIS STUDY?**

*State the* ***most important*** *reasons {benefits} a person may want to participate in this study.* For a complete Description of benefits please refer to the Detailed Consent.

**WHAT ARE THE REASONS YOU MIGHT NOT CHOOSE TO VOLUNTEER FOR THIS STUDY?**

*State the* ***most important*** *reasons {risks} a person may NOT want to participate in this study.* For a complete Description of risks please refer to the Detailed Consent.

**DO YOU HAVE TO TAKE PART IN THIS STUDY?**

You do not have to take part in this research. It is your choice whether or not you want to take part. You can agree to take part and later change your mind. If you choose not to take part or choose to stop taking part at any time, there will be no penalty to you or loss of benefits to which you are otherwise entitled.

*Add the following for student volunteers and/or employees:* As a student/employee, if you decide not to take part in this study, your choice will have no effect on your academic status or class grade(s) or employment status.

**WHAT IF YOU HAVE QUESTIONS OR CONCERNS?**

The person in charge of this study is \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ {Principal Investigator, PI}. If you have questions, suggestions, or concerns regarding this study or you want to withdraw from the study his/her contact information is: {PI contact information}.

This research is being overseen by an Institutional Review Board (“IRB”). You may talk to them at 202-994-2715 or via email at [ohrirb@gwu.edu](file:///C%3A%5CUsers%5Crescxr%5CAppData%5CLocal%5CTemp%5Cohrirb%40gwu.edu) if:

1. You have questions, concerns, or complaints that are not being answered by the research team or if you wish to talk to someone independent of the research team.
2. You have questions about your rights as a research subject.

## Detailed Consent Form:

## Why am I being invited to take part in a research study?

We invite you to take part in a research study because \_\_\_\_\_\_\_\_\_\_\_\_\_. [Fill in the circumstance or condition that makes subjects eligible for the research].

## What should I know about a research study?

1. Someone will explain this research study to you. You may ask all the questions you want before you decide whether to participate.
2. Participation is voluntary; whether or not you take part is up to you.
3. You can agree to take part and later change your mind.
4. Your decision not to take part or to stop your participation will not be held against you.
5. Your decision will not affect the medical care you receive from GW. If you decide not to take part, you can still receive medical care from GW.
6. You may take this document home to read or to discuss with your family members or doctor before deciding to take part in this research study.

*[Any applicable trials initiated on or after March 7, 2012, must include a specific statement that refers subjects to the publishing of the study data on www.ClinicalTrials.gov. The FDA provides the following language that must be used in all consent forms word-for-word]**ClinicalTrials.gov* is a website that provides information about federally and privately supported clinical trials. A description of this clinical trial will be available on [http://www.ClinicalTrials.gov](http://www.clinicaltrials.gov/), as required by U.S. Law. This website will not include information that can identify you. At most, the website will include a summary of the results. You can search this website at any time.

## Who can I talk to if I have questions?

If you have questions, concerns, or complaints, or think the research has hurt you, talk to the Principal Investigator at [Insert contact information for the PI]

This research is being overseen by an Institutional Review Board (“IRB”). You may talk to them at 202-994-2715 or via email at [ohrirb@gwu.edu](file:///C%3A%5CUsers%5Cresgxf%5CAppData%5CLocal%5CMicrosoft%5CWindows%5CTemporary%20Internet%20Files%5CContent.Outlook%5CHYC7ZE95%5Cohrirb%40gwu.edu) if:

1. You have questions, concerns, or complaints that are not being answered by the research team or if you wish to talk to someone independent of the research team.
2. You have questions about your rights as a research subject.

## Why is this research being done?

[Tell the subject the purpose of the research. Explain the background of the research problem. Explain any potential benefits to other. Describe any procedures that are important to the research that will be performed regardless of whether the subject takes part in the research.]

## How long will I be in the study?

We expect that you will be in this research study for \_\_\_\_\_\_\_\_[hours/days/months/weeks/years, until a certain event]**.**

## How many people will take part in this research study?

We expect about \_\_\_\_\_ people will take part in the entire study.

## What happens if I agree to be in this research?

[Tell the subject what to expect using lay language and simple terms. Include all procedures performed because the subject is taking part in the research, including procedures to monitor subjects for safety or minimize risks. Do NOT describe procedures that will be performed regardless of whether the subject takes part in the research. Describe these procedures in the section titled “Why is this research being done?

Whenever appropriate include the following items:

* A time-line description of the procedures that will be performed. If practical, prepare a time-line chart or schematic to accompany descriptions of procedures and tests for research that require more than 1 or 2 steps/visits
* The drugs or biologics that will be given to the subject
* All devices that will be used
* All hospitalizations, outpatient visits and telephone or written follow-up
* The length and duration of visits and procedures
* If blood will be drawn, indicate the amount [in English units] and frequency
* With whom will the subject interact
* Where the research will be done
* When the research will be done
* List experimental procedures and therapies and identify them as such
* How often procedures will be performed
* What is being performed as part of the research study
* What is being performed as part of standard care
* What procedures are part of regular medical care that will be done even if the subject does not take part in the research
* When applicable indicate that the subject will be contacted for future research.

[Include for a clinical trial that involves randomization. Otherwise delete]. The treatment you get will be chosen by chance, like flipping a coin. Neither you nor the study doctor will choose what treatment you get. You will have an \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ [equal/one in three/etc]. chance of being given each treatment. [For double-blinded research add]Neither you nor the people conducting the research will know which treatment you are getting. [For single blinded research add] You will not be told which treatment you are getting, however the people conducting the research will know.

## What are my responsibilities if I take part in this research?

[Delete this section if the research is not a clinical trial].

If you take part in this research, you will be responsible to: [Describe any responsibilities of the subject].

## What other choices do I have besides taking part in the research?

[Include if there are alternatives other than participating. Otherwise delete].Instead of being in this research study, your choices may include: [List alternatives procedures. For student subject pools describe alternatives for course credit. For clinical trials describe the options that you would normally offer patient. If applicable, include supportive care as an option. If drug/device/treatment is available as standard of care, state so here].

## What happens if I agree to be in research, but later change my mind?

[Include for research involving students or employees. Otherwise delete].If you are a student or employee at George Washington University/George Washington University Hospital or the MFA, your academic standing/employment status will not be affected in any way should you choose not to take part or to withdraw at any time.

[Include for all research]You may refuse to participate or you may discontinue your participation at any time without penalty or loss of benefits to which you would otherwise be entitled.

[Include if there are potential adverse medical consequences to withdrawing from the research. Otherwise delete] If you decide to leave the research, [Describe the adverse consequences].

If you decide to leave the research, contact the investigator so that the investigator can [Describe the procedures for orderly termination by the subject, if any].

[Include for FDA-regulated research. Otherwise delete]. If you stop being in the research, already collected data may not be removed from the study database. You will be asked whether the investigator can collect data from your routine medical care. *[Note: The consent document cannot give the subject the option of having data removed].*If you agree, this data will be handled the same as research data. *[Note: If a subject withdraws from the interventional portion of a study and does not consent to continued follow-up of associated clinical outcome information, the investigator must not access for purposes related to the study the subject’s medical record or other confidential records requiring the subject’s consent. However, an investigator may review study data related to the subject collected prior to the subject’s withdrawal from the study, and may consult public records, such as those establishing survival status].*

*[For research that is not FDA-regulated, describe what will happen to data collected to the point of withdrawal. Describe whether subjects will be asked to explain the extent of their withdrawal and whether they will be asked for permission to collect data through interaction or collection of private identifiable information. For example, a subject may wish to withdraw from the experimental procedure because of unacceptable side effects, but may agree to undergo follow-up procedures and data collection].*

## Is there any way being in this study could be bad for me?

 [Include for research that involves only minimal risks, otherwise delete]. The risks and discomforts associated with participation in this study are not greater than those ordinarily encountered in daily life or during the performance or routine physical or psychological examinations or tests.

[The risks of procedures may be presented in a table form].

[Describe each of the following risks, if appropriate. If known, describe the probability and magnitude of the risk].

* [Physical risks
* Psychological risks
* Privacy or confidentiality risks
* Legal risks
* Social risks
* Economic risks]

[Include for research that involves procedures whose risk profile is not well known, including all research involving an investigational product. Otherwise delete]. In addition to these risks, this research may hurt you in unknown ways. These may be a minor or so severe as to cause death.

[Include for research that involves pregnant women or women of child-bearing potential and procedures that involve risks to an embryo or fetus or whose risk profile in pregnancy is not well known. Otherwise delete].The procedures in this research can hurt a pregnancy or fetus in the following ways: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_. [Omit the previous sentence if there are no known risks]. The research may hurt a pregnancy or fetus in unknown ways. These may be a minor or so severe as to cause death. [Omit the previous two sentences for research whose risk profile in pregnancy is well known]. You should not be or become pregnant [include as applicable “*or father a baby”]*while on this research study.

***[***Include for research that *may result in additional costs to the subjects. Otherwise delete].* Taking part in this research study may lead to added costs to you. [Describe what these costs are].

[Include for a clinical trial. Otherwise delete]. You and your insurance company will be charged for the health care services that you would ordinarily be responsible to pay. In some cases, insurance will not pay for services ordinarily covered because these services were performed in a research study. You should check with your insurance to see what services will be covered by your insurance and what you will be responsible to pay.

[Include for a clinical trial where subjects are randomized to 2 standard of care procedures. Otherwise delete]. The risks and potential complications are similar if you decide to not participate in the study. One potential risk of participating in the study is that you may receive a different *[procedure, treatment, drug, device]* than you would if you do not participate. While these *[procedures, treatments, drugs, devices]* are currently believed to be safe, there may be additional risks discovered in the future. Your care will be identical whether you decide to participate in this study or not. If you do not participate in the study, the decision to have [procedure, treatment, drug, device] is up to the personal preference of you and your doctor.

**What happens if I believe I am injured because I took part in this study?**

The researchers have taken steps to minimize the known or expected risks. In spite of all precautions, you still may experience medical complications or side effects from participating in this study. You should promptly notify the study doctor in the event of any illness or injury as a result of being in the study.

[Include if compensation for research related injury is not available. Otherwise delete and insert language detailing compensation and medical treatment available.] If you believe that you have been injured or have become ill from taking part in this study, you should seek medical treatment from GWU Hospital and/or the GWU MFA or through your physician or treatment center of choice. Care for such injuries will be billed in the ordinary manner to you or your insurance company.

You will not receive any financial payments from GWU, GWU Hospital and/or the GWU MFA for any injuries or illnesses. You do not waive any liability rights for personal injury by signing this form.

## Will being in this study help me in any way?

[Include if there are no benefits to participation. Otherwise delete]. You will not receive any benefits from participating in this research.

[Include if there are benefits to participation. Otherwise delete]. We cannot promise any benefits to you or others from your taking part in this research. However, possible benefits include \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_. [Then describe the potential benefits of participation. First describe any direct benefits to the subject, then any benefits to others. If benefits from participation may not continue after the research has ended, describe them here. Monetary reimbursement for participation is not a benefit].

[Include for a clinical trial with no benefits to participation. Otherwise delete]. There are no benefits to you from your taking part in this research. We cannot promise any benefits to others from your taking part in this research. However, possible benefits to others include \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_. [Describe any benefits to others. Monetary reimbursement for participation is not a benefit and should be described in a later section].

[Include for research involving prisoners] Taking part in this research study will not improve your housing or correctional program assignments. Your taking part in this research study will not improve your chance of parole or release.

## Can I be removed from the research without my permission?

[Delete this section if not applicable].

[Include for research where this is a possibility. Otherwise delete]. The person in charge of the research study or the sponsor can remove you from the research study without your approval. Possible reasons for removal include [describe reasons why the subject may be withdrawn, if appropriate].

[Include for research where this is a possibility. Otherwise delete]. We will tell you about any new information that may affect your health, welfare, or choice to stay in the research.

## What happens to my information collected for the research?

To the extent allowed by law, we limit your personal information to people who have to review it. We cannot promise complete secrecy. The IRB and other representatives of this organization may inspect and copy your information. Others include [Add to this list other organizations that may have access to the subject’s records such as the Food and Drug Administration, when the research if FDA-regulated, the Department of Health and Human Services, when the research is conducted or funded by DHHS, the sponsor, contract research organization, sponsor’s agent and other collaborating institutions].

[Describe any limitations on confidentiality based on possible legal issues. For example, if the research team is likely to uncover abuse, neglect, or reportable diseases, explain that this information may be disclosed to appropriate authorities].

[Include if requesting a Certificate of Confidentiality. Otherwise delete]. To help us protect your privacy, we have obtained a Certificate of Confidentiality from the National Institutes of Health. The researchers can use this Certificate to legally refuse to disclose information that may identify you in any federal, state, or local civil, criminal, administrative, legislative, or other proceedings, for example, if there is a court subpoena. The researchers will use the Certificate to resist any demands for information that would identify you, *[except as explained below]*. *[Use the following language as applicable]* The Certificate cannot be used to resist a demand for information from personnel of the United States Government that is used for auditing or evaluation of Federally funded projects or for information that must be disclosed in order to meet the requirements of the federal Food and Drug Administration (FDA). You should understand that a Certificate of Confidentiality does not prevent you or a member of your family from voluntarily releasing information about yourself or your involvement in this research. If an insurer, medical care provider, or other person obtains your written consent to receive research information, then the researchers will not use the Certificate to withhold that information. *[Language such as the following should be included if researcher intend to make voluntary disclosure about information obtained in the research such as child abuse, or intent to hurt self or others.]* The Certificate of Confidentiality will not be used to prevent disclosure to state or local authorities of *[list what will be reported, such as child abuse and neglect, or harm to self or others]*.

 [Include for a clinical trial. Otherwise delete]. The sponsor, monitors, auditors, the IRB, the Food and Drug Administration will be granted direct access to your medical records to conduct and oversee the research. By signing this document you are authorizing this access. We may publish the results of this research. However, we will keep your name and other identifying information confidential.

*[The following language is recommended when samples of tissues, cells, blood, or body fluids (hereafter referred to as tissues) will be taken or banked for use in current or future research. This includes testing the sample for purposes of collecting genetic or other information. Investigators should choose the appropriate provisions to be included in their informed consent form and may vary any of the following language as appropriate.]*

Tissue Sampling for Research

Research using tissues is an important way to try to understand human disease. You have been given this information because the investigators want to include your tissues in a research project and because they want to save the samples for future research. There are several things you should know before allowing your tissues to be studied.

Your tissues will be stored *[insert how samples will be stored - and if appropriate how samples will be linked) e.g., under diagnosis and medical record or code number and unlinked]****.***

*[If linked]:* You have the right to refuse to allow your tissues to be studied now or saved for future study. You may withdraw from this study at any time. The investigators might retain the identified samples, e.g., as part of your routine clinical care, but not for additional research. De-identified information or biospecimens may or may not be used or shared for future research.

*[If unlinked]:* Because your samples will not be linked to your name after they are stored, you cannot withdraw your consent to the use of the samples after they are taken.

*[Optional]:*

The results of the study of your samples will be used for research purposes only and you will not be told the results of the tests.

*[Or]*

You will be told the results of tests that are part of your clinical care, but you will not be told the results of the research tests, including any future research tests.

 \_\_\_\_\_ I consent to my samples being saved for future research

 \_\_\_\_\_ I do not consent to my samples being saved for future research

Tissue Sampling for Genetic Testing

*[Include the following language if samples in study will be used for genetic testing or if future research on samples will include genetic testing.]*

As part of the analysis on your samples, the investigators *(may/will)* do genetic testing. Genetic research is research that studies genes, including gene characteristics and gene versions that are transmitted by parents to children. Genetic research may include looking at information, such as personal appearance and biochemistry, gene sequences, genetic landmarks, individual and family medical histories, reactions to medications and responses to treatment. Genetic research raises certain questions about informing you of any results. Possible risks of knowing results include: anxiety; other psychological distress; and the possibility of insurance and job discrimination. A possible risk of not knowing includes being unaware of the need for treatment. These risks can change depending on the results of the research and whether there is a treatment or cure for a particular disease.

Sometimes patients have been required to furnish information from genetic testing for health insurance, life insurance, and/or a job. A Federal law, the Genetic Information Nondiscrimination Act of 2008 (GINA), generally makes it illegal for health insurance companies, group health plans, and employers with 15 or more employees to discriminate against you based on your genetic information.

*[If investigators will not share the research results with the participant, the following language can be added]:*

The results of the study of your samples from this project will be used for research purposes only, and you will not be told the results of the tests.

*[If investigators will allow participants to choose whether they want to receive test results and/or will contact participants in the future, the following language (two choices of language) can be added]:*

Regarding informing you of the test results, you should understand the following:

* The information may be too limited to give you particular details or consequences;
* You may be determined to carry a gene for a particular disease that can be treated;
* You may be determined to carry a gene for a particular disease for which there is no current treatment;
* You carry a gene for a disease and might consider informing relatives that they, too, might carry the gene.

Please circle [ yes or no ] as to whether you wish to be told the test results.

Please circle [ yes or no ] as to whether you wish your family members to be told the test results.

*[Or]*

Investigators in this study may try to re-contact you in the future. If you are re-contacted and want to know what the investigators have learned about your tissue samples, you should understand the following:

* The information may be too limited to give you particular details or consequences;
* You may be determined to carry a gene for a particular disease that can be treated;
* You may be determined to carry a gene for a particular disease for which there is no current treatment;
* You carry a gene for a disease and might consider informing relatives that they, too, might carry the gene.

*[If you would like to contact participants about future studies, include the following statement]:*

**\***May we contact you about future studies that may be of interest to you?

\_\_\_\_ Yes \_\_\_\_ No

How will my privacy and health information be protected?

The Health Insurance Portability and Accountability Act (HIPAA) requires that researchers and health care providers protect the privacy of information that identifies you and relates to your past, present and future physical and mental health or conditions, or the provision of health care. If you agree to participate in this research, protected health information will be used and shared with others for purposes of the study. Below is more detailed information about how your health information will be shared and protected. By signing this form, you are allowing the people and groups that are listed in the next paragraphs to use your health information for this research study. Your information will only be used or shared as explained in this authorization form

The use and release of protected health information is for the purpose of collecting data for this study.

Protected Health Information to be shared: [list all PHI that will be used or disclosed for this specific study]

Who may disclose your protected health information: The researcher and the other members of the research team may obtain your individual health information from:

Hospitals: <List by name>

Clinics: <List by name>

Other Providers: <List by name>

Health Plan: <List by name>

And from hospitals, clinics, health care providers, and health plans that provide health care to you during the study [NOTE: Delete types of CE’s that do NOT apply to this study]

By signing this form, you allow the use, sharing, copying, and release of your protected health information in connection with this study by:

• The members of the research team;

• Other healthcare providers such as labs which are part of the study;

• A safety monitoring board {include only if applicable};

• Institutional officials who are responsible for compliance;

Some of the tests in this study would have been done as part of your regular care. These test results will be used both to treat you and to complete this research. The test results will be recorded in your medical record. These study results will be included in your medical record. Results of tests and studies done solely for this research study and not as part of your regular care will [also] OR [not] be included in your medical record.

[or use this alternative statement:]

All tests are being done only because you are in this study. The study results will [not] be [given to you to send] OR [sent] to your physician to include in your medical record

Once your health information has been disclosed to others outside of the hospitals and medical practices [customize this part of the phrase to fit this study], the information may no longer be covered by the federal regulation that protects privacy of health information.

Not signing this form or later canceling your permission will not affect your health care treatment outside the study, payment for health care from a health plan, or ability to get health plan benefits. However, if you do not give permission to use your health information, you may not take part in this study because your health information is needed in order to conduct this study.

This Authorization does not have an expiration date.

However, you may cancel this authorization at any time. Even if you cancel this authorization, the researchers may still use the protected health information they already have about you; however, no new health information or new biological specimens will be collected from you after you cancel your permission.

To cancel your permission, you will need to send a letter to [name Principal Investigator] stating that you are canceling your authorization. This letter must be signed and dated and sent to this address: [enter the name and address of the Principal Investigator].

 [Include for research involving prisoners. Otherwise delete]. If you are a prisoner, your medical records may also be given to officials and agencies within the criminal justice system when necessary and permitted by law.

## Are there any costs for participating in this research?

*[Include any costs that may be incurred due to the research. Indicate whether participants will be financially responsible for any clinic/hospital charges.]*

## Will I be paid for my participation in this research?

[Include if subjects will be paid. Otherwise delete].If you agree to take part in this research study, we will pay you \_\_\_\_\_\_\_\_ [indicate amount] for your time and effort. [Indicate if the amount is pro-rated for research visit completion].

[Include if subjects will be paid $600 or more during the course of the study. Otherwise delete.] Your name and social security number will be reported to the appropriate George Washington University (or GW Medical Faculty Associate) employees for purposes of making and recording the payment. You are responsible for paying income taxes on any payments provided by the study. If you receive $600 or more from the University (or the GW Medical Faculty Associate), GW must report the amount you receive to the Internal Revenue Service (IRS) on the form 1099-Misc. This form tells the IRS that payment was made to you, but it does not say that you were paid for taking part in this research study. You should talk to your tax advisor regarding the proper use of this form 1099-Misc.

If the subject’s biospecimens (even if identifiers are removed) may be used for commercial profit, include the following statement: (Modify if subjects will share in commercial profit.)

Your specimens (even if identifiers are removed) may be used for commercial profit. You will not share in this commercial profit.

## What else do I need to know?

[Include for sponsored research. Otherwise delete].This research is being funded by [Insert name of sponsor]**.**

[Include GINA language if conducting genetic testing] A federal law, called the Genetic Information Nondiscrimination Act (GINA), generally makes it illegal for health insurance companies, group health plans, and most employers to discriminate against you based on your genetic information. This law generally offers the following protections:

• Health insurance companies and employer-based group health plans may not request your genetic information that we get from this research.

• Health insurance companies and employer-based group health plans may not use your genetic information when making decisions regarding your eligibility or premiums.

• Employers with 15 or more employees may not use your genetic information that we get from this research when making a decision to hire, promote, or fire you or when setting the terms of your employment.

• All health insurance companies and group health plans must follow this law. All employers with 15 or more employees must follow this law as of November 21, 2009. The protections offered by GINA apply regardless of when the research that obtained the genetic information was conducted, even if prior to the effective date.

Be aware that this law does not protect you against discrimination on the basis of your genetic information by companies that sell life insurance, disability insurance, or long-term care insurance.

*[Include when applicable a statement indicating that the research will or might include whole genome sequencing]*

[Include for research involving prisoners where there may be a need for follow-up examination or care after the end of participation. Otherwise delete]. If you are released from jail before you finish this research study, you should take steps to get insurance or Medicaid coverage. Regular office visits and standard treatment will be billed to you or your health insurance. You may continue in the research study after your release from prison. If you move out of the area, we will help you make arrangements to be followed by a physician.

*[When applicable indicate that the investigator believes that the biologic specimens obtained could be part of or lead to the development of a commercial product].*

*[When applicable indicate when and how the subject will be informed of the results of the research. Include a statement that indicates whether clinical results, including individual research results, will be returned to the subject, and if so, under what conditions].*

[There are three signature pages attached to this template consent. Use the signature page or pages appropriate for your study. The IRB recommends that you make separate consent documents for each signature page to be used].

[Omit the signature page if there is no written documentation of consent].

[Include if investigators have a conflict of interest, and tailor to specific conflict. Otherwise delete.] An individual responsible for the conduct of this research study has a financial interest in the study drug or the medical device being used in this research study.

**Signature Block for Adult**

By signing below, you agree that the above information has been explained to you and you have had the opportunity to ask questions. You understand that you may ask questions about any aspect of this research during the course of the study and in the future. Your signature documents your permission to take part in this research. Your signature documents your permission to take part in this research.

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Printed name of subject

 \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_

Signature of subject Date

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_

Signature of person obtaining consent Date

 *[Add the following block if a witness will observe the consent process. E.g., short form of consent documentation or illiterate subjects].*

My signature below documents that the information in the consent document and any other written information was accurately explained to, and apparently understood by, the subject, and that consent was freely given by the subject.

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_

Signature of witness to consent process Date