Title of research study: [insert title of research study here with protocol number, if applicable]

[PLEASE BE SURE TO INSERT INFORMATION BELOW. List principal investigator(s), sub-investigators, sponsors and investigational sites where facilities will be used.]

Example:

Principal Investigator(s): XXXXX, Ph.D.

Sub-Investigator(s): XXXXXX, PhD
XXXXXX, MA
XXXXXX, MD

Faculty Supervisor: [If principal investigator is a graduate student, otherwise delete]

Sponsor: [Company/Federal Sponsor/UCF Department]
[Delete if study is not funded]

Investigational Site(s): [e.g. Orlando Regional Medical Center, Winter Park High School, University of Central Florida, Department...etc.]
Permission to Take Part in a Human Research Study

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?
- Someone will explain this research study to you.
- Whether or not you take part is up to you.
- You can choose not to take part.
- You can agree to take part and later change your mind.
- Your decision will not be held against you.
- You can ask all the questions you want before you decide.

Who can I talk to?
If you have questions, concerns, or complaints, or think the research has hurt you, talk to the research team at [Insert contact information for the research team]

This research has been reviewed and approved by an Institutional Review Board (“IRB”). You may talk to them at 407-823-2901 or irb@mail.ucf.edu.org if:
- Your questions, concerns, or complaints are not being answered by the research team.
- You cannot reach the research team.
- You want to talk to someone besides the research team.
- You have questions about your rights as a research subject.
- You want to get information or provide input about this research.

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 others.]

How long will the research last?
We expect that you will be in this research study for _______ [hours/days/months/weeks/years, until a certain event].

How many people will be studied?
We expect about _____ people here will be in this research study out of _____ people in the entire study nationally [or internationally].

What happens if I say yes, I want to be in this research?
[Tell the subject what to expect using lay language and simple terms. 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
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- 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 study doctor will know which treatment you are getting. [For single blinded research add] You will not be told which treatment you are getting, however your study doctor 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 happens if I do not want to be in this research?

Participation in research is completely voluntary. You can decide to participate or not to participate.

[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.]

[Include for a clinical trial. Otherwise delete.] The important risks and possible benefits of these alternatives include: [Describe the important risks and potential benefits of the alternative procedures and courses of treatment.]

What happens if I say yes, but I change my mind later?

You can leave the research at any time it will not be held against you.

[Include if there are potential adverse 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]
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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?

[Delete this section if there are no risks or discomforts.]

[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 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 ways that are unknown. These may be a minor inconvenience or may be 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 are known to hurt a pregnancy or fetus in the following ways: _______________. [Omit the previous sentence if there are no known risks.] The research may also hurt a pregnancy or fetus in ways that are unknown. These may be a minor inconvenience or may be 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.

Will being in this study help me any way?

[Delete this section if there are no benefits and the research is not a clinical trial.]

[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.

What happens to the information collected for the research?

Efforts will be made to limit the use and disclosure of your personal information, including research study and medical records, to people who have a need to review this information. We cannot promise complete secrecy. Organizations that may inspect and copy your information include the IRB and other representatives of this organization. [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.]

[If data or specimens will be retained after the study for future research, explain where the data or specimens will be stored, who will have access to the data or specimens, and how long the date or specimens will be retained.]

[Include for research where the sponsor may pay for medical expenses of the subject.] If the sponsor pays any of your medical expenses, we may be required to give the sponsor your name, date of birth, and Medicare ID or social security number.

[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.

[Include for FDA-regulated controlled drug and device trials (except Phase I drug trials) and FDA-regulated pediatric post-market surveillance trials of devices. Otherwise delete.] A description of this clinical trial will be available on http://www.ClinicalTrials.gov, as required by U.S. Law. This Web site will not include information that can identify you. At most, the Web site will include a summary of the results. You can search this Web site at any time.

[Include if a HIPAA authorization is required. Otherwise delete.] Federal law provides additional protections of your medical records and related health information. These are described in an attached document.

[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.

Can I be removed from the research without my OK?

[Delete this section if not applicable.]
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[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 else do I need to know?
[Include for sponsored research. Otherwise delete.] This research is being funded by [Insert name of sponsor].

[Include for research involving more than minimal risk. Otherwise delete.] If you need medical care because of taking part in this research study, contact the investigator and medical care will be made available. Generally, this care will be billed to you, your insurance, or other third party. [Insert the name of the institution] has no program to pay for medical care for research-related injury. [Describe any compensation available for research related injury.]

[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 for Department of Defense (DOD) research that targets military personnel where subjects will be paid. Otherwise delete.] Military personnel should check with their supervisor before accepting payment for participation in this research.

[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.]

[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.]
Permission to Take Part in a Human Research Study

Signature Block for Capable Adult

Your signature documents your permission to take part in this research.

Signature of subject

Date

Printed name of subject

Signature of person obtaining consent

Date

Printed name of person obtaining consent

IRB Approval 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

Printed name of person witnessing consent process
Permission to Take Part in a Human Research Study

Signature Block for Adult Unable to Consent
Your signature documents your permission for the named subject to take part in this research.

__________
Printed name of subject

__________
Signature of legally authorized representative

__________
Printed name of legally authorized representative

__________
Signature of person obtaining consent

__________
Signature of person obtaining consent

[Add the following block if you will document assent of the subject.]

☐ Obtained
☐ Not obtained because the capability of the subject is so limited that the subject cannot reasonably be consulted.

[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

__________
Printed name of person witnessing consent process
Permission to Take Part in a Human Research Study

Signature Block for Children

Your signature documents your permission for the named child to take part in this research.

Printed name of child

Signature of parent or individual legally authorized to consent to the child’s general medical care

Date

☐ Parent

☐ Individual legally authorized to consent to the child’s general medical care (See note below)

Note: Investigators are to ensure that individuals who are not parents can demonstrate their legal authority to consent to the child’s general medical care. Contact legal counsel if any questions arise.

Signature of parent

Date

Printed name of parent

If signature of second parent not obtained, indicate why: (select one)

☐ The IRB determined that the permission of one parent is sufficient. [Delete if the IRB did not make this determination]

☐ Second parent is deceased

☐ Second parent is unknown

☐ Second parent is incompetent

☐ Second parent is not reasonably available

☐ Only one parent has legal responsibility for the care and custody of the child

[Add the following block if you will document assent of children]

Assent

☐ Obtained

☐ Not obtained because the capability of the child is so limited that the child cannot reasonably be consulted.

[Add the following block to all consents]

Signature of person obtaining consent and assent

Date

Printed name of person obtaining consent

IRB Approval 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

Printed name of person witnessing consent process

Document Revision Date: December 12, 2017