University of Idaho
Permission to Use Personal Health Information for Research

Study Title (or IRB Approval Number if study title may breach subject's privacy):
_____

Sponsor/Funding Agency (if funded):
_____

A. What is the purpose of this form?

State and federal privacy laws protect the use and release of your health information. Under these laws, the University of Idaho or your health care provider cannot release your health information to the research team unless you give your permission. The research team includes the researchers and people hired by the University or the sponsor to do the research. If you decide to give your permission and to participate in the study, you must sign this form as well as the Consent Form. This form describes the different ways that the researcher, research team and research sponsor may use your health information for the research study. The research team will use and protect your information as described in the attached Consent Form. However, once your health information is released it may not be protected by the privacy laws and might be shared with others.

B. What Personal Health Information will be released?

If you give your permission and sign this form, you are allowing [Healthcare Provider] to release the following medical records containing your Personal Health Information. Your Personal Health Information includes health information in your medical records and information that can identify you. For example, Personal Health Information may include your name, address, phone number or social security number.

☐ Entire Medical Record
☐ Radiology Reports
☐Pathology Reports
☐ Laboratory Reports
☐ Dental Records
☐ Operative Reports
☐ Emergency Medicine Center Reports
☐ Progress Notes
☐ History & Physical Exams
☐ Discharge Summary
☐ Consultations
☐ Outpatient Clinic Records
☐ EKG
☐ Radiology images
☐ Psychological Tests
☐ Health Care Billing Statements

☐ Other:
_____

C. Do I have to give my permission for certain specific uses?

Yes. The following information will only be released if you give your specific permission by putting your initials on the line(s).

____ I agree to the release of information pertaining to drug and alcohol abuse, diagnosis or treatment.
____ I agree to the release of HIV/AIDS testing information.
____ I agree to the release of genetic testing information.
____ I agree to the release of information pertaining to mental health diagnosis or treatment as follows: _____.

D. How will my Personal Health Information be used?

Your Personal Health Information may be released to these people for the following purposes:

1. To the research team for the research described in the Consent Form;
2. To others at University who are required by law to review the research;
3. To others who are required by law to review the quality and safety of the research, including: U.S. government agencies, such as the Food and Drug Administration, the research sponsor or the sponsor's representatives, or government agencies in other countries. These organizations and their representatives may see your Personal Health Information. They may not copy or take it from your medical records unless permitted or required by law.

E. How will my Personal Health Information be used in a research report?

If you agree to be in this study, the research team may fill out a research report. (This is sometimes called "a case report".) The research report will **not** include your name, address, or telephone or social security number. The research report may include your date of birth, initials, dates you received medical care, and a tracking code. The research report will also include information the research team collects for the study. The research team and the research sponsor may use the research report and share it with others in the following ways:

1. To perform more research;
2. Share it with researchers in the U.S. or other countries;
3. Place it into research databases;
4. Use it to improve the design of future studies;
5. Use it to publish articles or for presentations to other researchers;
6. Share it with business partners of the sponsor; or
7. File applications with U.S. or foreign government agencies to get approval for new drugs or health care products.

F. Does my permission expire?

This permission to release your Personal Health Information expires when the research ends and all required study monitoring is over. Research reports can be used forever.

G. Can I cancel my permission?

You can cancel your permission at any time by writing to the researcher to inform him or her that you wish to cancel your permission. If you cancel your permission, you may no longer be in the research study. If you cancel, information that was already collected and disclosed about you may continue to be used. Also, if the law requires it, the sponsor and government agencies may continue to look at your medical records to review the quality or safety of the study.

H. Signature

If you agree to the use and release of your Personal Health Information, please sign below. You will be given a signed copy of this form**.**

_____

Subject's Name (print)

_____ _____

Subject's Signature Date

Note: if the subject is a minor, an individual signing with an "X", an adult incapable of giving consent, or is unable to read the authorization, fill out and attach the "special signatures" page (sections "I" and "J").

SPECIAL SIGNATURES PAGE

I. If the subject is a minor, or an individual signing with an "X", or an adult incapable of giving consent (where IRB approved), the legally authorized representative or witness signs here:

_____ _____
Legally Authorized Representative's Name or Witness to the "X" (print) | Relationship to the Subject

_____ _____
Representative or Witness Signature | Date

J. If the subject is unable to read the authorization, the translator or reader and a witness sign here:

I have accurately and completely read this Authorization to _____ (subject's name) in _____(language), the subject's primary language. **The subject has verbally affirmed his/her Authorization to me and to the witness.**

_____
Translator or Reader's Name (print)

_____ _____
Translator or Reader's Signature | Date

_____
Witness Name (print)

_____ _____
Witness Signature | Date