<u>University of California</u> <u>Permission to Use Personal Health Information for Research</u>

Study Title (or IRB Approval Number if study title may breach subject's privacy):				
Sponsor/Funding Agency (if funded):				
laws, the University of Californ the research team unless you people hired by the University permission and to participate form describes the different your health information for the information as described in the released it may not be protequestions, ask a member of the secondary of the sec	ws protect the use and release prints or your health care provided give your permission. The ty or the sponsor to do the rese in the study, you must sign the ways that the researcher, researcher attached Consent Form. It is the attached Consent Form. It is the research team. Information will be released and sign this form, you are allower(s) releasing medical records ealth Information. Your Person	owing [insert UC campus or state of the following medical records hall Health Information includes health an identify you. For example, Personal Health		
 □ Entire Medical Record □ Health Care Billing Statements □ Pathology Reports □ EKG 	□ Dental Records□ Operative Reports	 □ Emergency Medicine Center Reports □ History & Physical Exams □ Diagnostic Imaging Reports □ Consultations 		
☐ Progress Notes ☐ Other (describe)	□ Radiology Reports□ Radiologic & MR Scans□ Discharge Summary	 □ Outpatient Clinic Records □ Psychological Tests 		

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 attached Consent Form;
- 2. To others at UC 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.

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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. You can do this in two ways. You can write to the researcher or you can ask someone on the research team to give you a form to fill out to cancel your permission. If you cancel your permission, you may no longer be in the research study. You may want to ask someone on the research team if canceling will affect your medical treatment. 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 Persegiven a signed copy of this form.	onal Health Information, please sign below. You will be
Subject's Name (print)	
Subject's Signature	 Date

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H. If the subject is a minor, or an individual signiconsent (where IRB approved), the legally author	
Legally Authorized Representative's Name or Witness to the "X" (print)	Relationship to the Subject
Representative or Witness Signature	

RB Approval Number	
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H. If the subject is unable to read the authorizat here:	ion, the translator or reader and a witness sign
I have accurately and completely read this Authoriza name) in(language), the subject affirmed his/her Authorization to me and to the ways are also as a subject of the wa	's primary language. The subject has verbally
Translator or Reader's Name (print)	
Translator or Reader's Signature	 Date
Witness Name (print)	
Witness Signature	 Date