IRB#	
University of California	
(Name of Your Health System) ¹	
Permission to Use Personal Health Information for Research	
udy Title (or IRB Approval Number if study title may breach subject's privacy):	
incipal Investigator Name:	
oonsor/Funding Agency (if funded):	
What is the purpose of this form?	
ate and federal privacy laws protect the use and release of your health information of these laws, the University of California or your health care provider cannot lease your health information for research purposes unless you give your permission our information will be released to the research team which includes the research exple hired by the University or the sponsor to do the research and people with atthority to oversee the research. If you decide to give your permission and to articipate in the study, you must sign this form as well as the Consent Form. This forms of Health System Component) 2 can share your information with the researcher, research am, sponsor and people with oversight responsibility. The research team will use a otect your information as described in the attached Consent Form. However, once ur health information is released by	on. ers, orm ch
th others. If you have questions, ask a member of the research team.	
What Personal Health Information will be released?	
you give your permission and sign this form, you are allowing:	า

 ¹ Each UC Health System or business unit may elect to leave this as UC or add the name of their specific health system or unit.
² The name here should match how the organization is identified in the Notice of Privacy Practices.
³ The name here should match how the organization is identified in the Notice of Privacy Practices.

☐ Entire Medical Record	☐ Lab & Pathology Reports	☐ Emergency ☐ Records	Department
☐ Ambulatory Clinic Records	□ Dental Records	☐ Financial Red	cords
☐ Progress Notes	☐ Operative Reports	☐ Imaging Repo	orts
☐ Other Test Reports	☐ Discharge Summary	☐ History & Phy	
☐ Other (describe)	☐ Consultations	☐ Psychologica	
(Description of Other Health Information)			
C. Do I have to give my perm	ission for certain specific	uses?	
Yes. The following information by putting your initials on the I	-	give your specific	permission
I agree to the release of diagnosis or treatment.	information pertaining to dru	ıg and alcohol ab	use,
I agree to the release of	HIV/AIDS testing informatio	n.	
I agree to the release of	genetic testing information.		
	information pertaining to me	ental health diagn	osis or
D. Who will disclose and/or r	eceive my Personal Health	Information?	
Your Personal Health Informa purposes:	tion may be shared with thes	se people for the	following
_	r the research described in t hority to oversee the resear		sent Form;
including: U.S. governm	red by law to review the qua ent agencies, such as the Fo Research Protections, the re	ood and Drug Adı	•
			(Sponsor Name)
or the sponsor's represe	entatives including but not lin	nited to	
government agencies in	other countries		(CRO Name), OI
government agencies in	oner countiles.		

E. How will my Personal Health Information be shared for the research?

If you agree to be in this study, the research team may share your Personal Health Information in the following ways:

- 1. To perform the research
- 2. Share it with researchers in the U.S. or other countries;
- 3. Use it to improve the design of future studies;
- 4. Share it with business partners of the sponsor; or

5. File applications with U.S. or foreign government agencies to get approval for new drugs or health care products.

F. Am I required to sign this document?

No, you are not required to sign this document. You will receive the same clinical care if you do not sign this document. However, if you do not sign the document, you will not be able to participate in this research study.

G. Optional research activity

If the research I am agreeing to participate in has additional optional research activity such as the creation of a database, a tissue repository or other activities, as explained to me in the informed consent process, I understand I can choose to agree to have my information shared for those activities or not.

☐ I agree to allow my information to be disclosed for the additional optional research activities explained in the informed consent process.

H. Does my permission expire?

This permission to release your Personal Health Information expires when the research ends and all required study monitoring is over.

I. 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 for limited purposes. 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.

J. Signature

Subject	
If you agree to the use and release of your Perso your name and sign below. You will be given a sign	•
Subject's Name (print)—required	_
Subject's Signature	 Date

Parent or Legally Authorized Representative			
If you agree to the use and release of the above named subject's Personal Health Information, please print your name and sign below.			
Parent or Legally Authorized Representative's Nar (print)	me Relationship to Subject		
Parent or Legally Authorized Representative's Signature	Date		
Witness			
If this form is being read to the subject because s/witness must be present and is required to print his	•		
Witness' Name (print)			
Witness' Signature	Date		