Authorization for Use & Release of Individually Identifiable Health Information for Veterans Health Administration (VHA) Research			
Subject Name (Last, First, Middle Initial):	:	Subject SSN (last 4 only):	Date of Birth:
VA Facility (Name and Address):			
VA Principal Investigator (PI):		PI Contact Information:	
Study Title:			
Purpose of Study:			
USE OF YOUR INDIVIDUALLY IDENTIFIABLE HEALTH INFORMATION (IIHI): Your individually identifiable health information is information about you that contains your health information and information that would identify you such as your name, date of birth, or other individual identifiers. VHA is asking you to allow the VA Principal Investigator (PI) and /or the VA research team members to access and use your past or present health information in addition to new health information they may collect for the study named above. The investigators of this study are committed to protecting your privacy and the confidentiality of information related to your health care. Signing this authorization is completely voluntary. However, your authorization (permission) is necessary to participate in this study. Your treatment, payment, enrollment, or eligibility for VA benefits will not be affected,			
whether or not you sign this authorization. Your individually identifiable health information used for this VA study includes the information marked below: Information from your VA Health Records such as diagnoses, progress notes, medications, lab or radiology findings, etc.			
 □ Specific information concerning: □ alcohol abuse □ drug abuse □ sickle cell anemia □ HIV □ Demographic Information such as name, age, race, etc. □ Billing or Financial Records □ Photographs, Videotapes, and/or Audiotapes of you □ Questionnaire, Survey, and/or Subject Diary □ Other, as immediately described below: 			

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Subject Name (Last, First, Middle Initial):	Subject SSN (last 4 only):	Date of Birth:
USE OF YOUR DATA OR SPECIMENS FOR OTHER RESEARCH: banking is a required component of this study. When banking is an of this form in lieu of this section.)	•	-
	arch	
An important part of this research is to save your		
☐ Data		
☐ Specimen		
in a secure repository/bank for other research studies in the future. If and/or specimen for future studies approved by the required committe will not be able to participate in this study.	,	3
DISCLOSURE: The VA research team may need to disclose the info institutions that are not part of VA. VA/VHA complies with the require Accountability Act of 1996 (HIPAA), Privacy Act of 1974 and all other protect your privacy. The VHA Notice of Privacy Practices (a separate we protect your information. If you do not have a copy of the Notice, the Giving your permission by signing this authorization allows us to disclepersons outside the VA/VHA as noted below. Once your information longer be protected by federal laws and regulations and might be reducted information. These non-VA/VHA institutions or persons include the	ments of the Health Insurance applicable federal laws and redocument) provides more in research team will provide ose your information to other has been disclosed outside isclosed by the persons or in	e Portability and regulations that of the second se
Non-VA Institutional Review Board (IRB) at who will monitor the study		
Study Sponsor (name):		
Person or entity who takes responsibility for and initiates a clinical	investigation	
Academic Affiliate (institution/name/employee/department): A relationship with VA in the performance of this study		
Compliance and Safety Monitors:	A d	
Advises the Sponsor or PI regarding the continuing safety of this s		
Other Federal agencies required to monitor or oversee research (s	uch as FDA, OHRP, GAO):	
☐ A Non-Profit Corporation (name and specific purpose):		
☐ Other (e.g. name of contractor and specific purpose):		

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Subject Name (Last, First, Middle Initial):	Subject SSN (last 4 only):	Date of Birth:		
Note: Offices within VA/VHA that are responsible for oversight of VA research such as the Office of Research Oversight (ORO), the Office of Research and Development (ORD), the VA Office of Inspector General, the VA Office of General Counsel, the VA IRB and Research and Development Committee may also have access to your information in the performance of their VA/VHA job duties.				
Access to your Individually Identifiable Health Information created or obtained in the course of this research: While this study is being conducted, you				
☐ will have access to your research related health records				
☐ will not have access to your research related health records				
This will not affect your VA healthcare including your doctor's ability to see your records as part of your normal care and will not affect your right to have access to the research records after the study is completed.				
REVOCATION: If you sign this authorization you may change your mind and revoke or take back your permission at any time. You must do this in writing and must send your written request to the Principal Investigator for this study at the following address:				
If you revoke (take back) your permission, you will no longer be able to participate in this study but the benefits to which you are entitled will NOT be affected. If you revoke (take back) your permission, the research team may continue to use or disclose the information that it has already collected before you revoked (took back) your permission which the research team has relied upon for the research. Your written revocation is effective as soon as it is received by the study's Principal Investigator.				
EXPIRATION: Unless you revoke (take back) your permission, your authorization to allow us to use and/or disclose your information will:				
☐ Expire at the end of this research study				
☐ Expire on the following date or event:				
☐ Not expire				
Expires at the end of this research study unless you have: (1) provided additional permission to store your data and/or biological specimens in a research data repository or (2)when further optional analysis of your specimens has been completed				

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Authorization for Use & Release of Individually Identifiable Health Information for Veterans Health Administration (VHA) Research			
Subject Name (Last, First, Middle Initial):	Subject SSN (last 4 only):	Date of Birth:	
TO BE FILLED OUT BY THE SUBJECT			
Research Subject Signature. This permission (authorization) has be opportunity to ask questions. If I believe that my privacy rights have facility Privacy Officer to file a verbal or written complaint.			
I give my authorization (permission) for the use and disclosure of my described in this form. I will be given a signed copy of this form for m		h information as	
Signature of Research Subject	Date		
Signature of Legal Representative (if applicable)	Date		
To Sign for Research Subject (Attach authority to sign: Health Care P or Next of Kin if authorized by State Law)	ower of Attorney, Legal Gua	ardian appointment,	
Name of Legal Representative (please print)	Date		

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Authorization for Use & Release of Individually Identifiable Health Information for Veterans Health Administration (VHA) Research		
Subject Name (Last, First, Middle Initial):		
Subject Social Security Number (last 4 numbers only):	Date of Birth:	
VA Facility (Name and Address):		
VA Principal Investigator (PI):	PI Contact Information:	
Study Title:	I	
Optional Authorization Supplement for Placing My Data of Conducting Optional Analysis of My Specimens For Futu	or My Biological Specimens in a Repository or for are Use by Veterans Health Administration (VHA) Research	
Purpose. This supplement to the authorization is for eithe example blood, urine, tissue) collected during the study fo study. You are not required to provide this permission and participation in this study, i.e., granting this permission is n	r future research or for conducting optional analysis for this I not providing this permission will have no impact on your	
Research Subject Signature. This additional permission given the opportunity to ask questions about this activity.		
☐ Store my health information in a research data reposito	ry,	
☐ Store my biological specimens (blood, tissue, urine, etc	c.) in a research data repository, or	
☐ Further optional analysis of my specimens occurring be	elow:	
Future research of data maintained within a research data Board and/or other applicable approvals to ensure the prot	·	
Signature of Research Subject	Date	
Signature of Legal Representative (if applicable)	Date	
Name of Legal Representative (please print)	Date	
To Sign for Research Subject (Attach authority to sign: He or Next of Kin if authorized by State law)	ealth Care Power of Attorney, Legal Guardian appointment,	
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