

Request for a Full Waiver of Authorization Requirement

Sponsor		S	Sponsor Protocol No			
pers	on responsible for the con	duct of the research	ΓΙΟΝ: Please provide information about the h. HPH must be assured that the investigator h and the protection of human subjects.			
1.	PI Name:					
2.	PI Company Name:					
3.	PI Mailing Address: (street, city, state/province, zip, country)					
4.	PI Phone:	PI Fax:	PI E-mail:			
5.	How would the PI prefer to receive study documents? (check one) Fax E-mail Regular Mail					
Wai	Specify which, if any, of the following identifiers will be associated with the health information you propose to access for the study. Names Telephone Numbers					
	Address		E-mail Addresses			
	Fax Numbers		Medical Record Numbers			
	Social Security Numbers Health Plan Beneficiary Number Certificate/License Numbers		Account Numbers Vehicle Identifiers and Serial Numbers			
			Web Universal Resource Locators (URL)			
	Device Identifiers and Serial Numbers		Biometric Identifiers (finger and voice prints)			
	Internet Protocol (IP) Address Numbers					
	Any Geographic Subd		Any Elements of Dates (specify which of the			
	Than a State (specify which of the		following identifiers you will use: birth date,			
	following identifiers y		admission date, discharge date, date of death,			
	city, parish, or zip cod		age over 89):			
	Full face photographic comparable images:	images and	Any other unique identifying number, characteristic, or code (please specify):			

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2.		s will be associated with the health information you						
	propose to collect and use for the study.							
	Names	Telephone Numbers						
	Address	E-mail Addresses						
	Fax Numbers	Medical Record Numbers						
	Social Security Numbers	Account Numbers						
	Health Plan Beneficiary Number	Vehicle Identifiers and Serial Numbers						
	Certificate/License Numbers Device Identifiers and Serial Numbers	Web Universal Resource Locators (URL) Biometric Identifiers (finger and voice prints)						
	Internet Protocol (IP) Address Numbers							
	Any Geographic Subdivisions Smaller	Any Elements of Dates (specify which of the						
	Than a State (specify which of the	following identifiers you will use: birth date,						
	following identifiers you will use: county,	admission date, discharge date, date of death,						
	city, parish, or zip code):	age over 89):						
	Full face photographic images and	Any other unique identifying number,						
	comparable images:	characteristic, or code (please specify):						
	a computation magger.	characteristic, or come (produce species).						
3.	List the specific health information that you pro	opose to collect and use in this study. State						
	specifically whether sensitive information (e.g., illegal drug use, sexual practices, HIV status) will be							
	collected. For most retrospective medical record	research, a limited range of health information will						
	normally be sufficient for the purposes of the research. A copy of the data collection sheet also							
	should be submitted for medical record review or database research studies. For survey or interview							
	research, the questions to be asked of research su	bjects should be attached to this application.						
		. ((2777))						
4.	4. What is the source of the Protected Health Information ("PHI")? List all sources from which you							
	plan to obtain PHI for the study (e.g. Facility or clinic paper records, a departmental database, your							
	own database)							
5.	Who will have access to, receive and/or use the information?							
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6.	Are the persons who have access to the information required to sign confidentiality Yes N							
	statements?							
7.	What identifiers are included on the information you plan to disclose?							
8.	List, if any, the individuals or groups outside of HPH to whom you will disclose the PHI (e.g.,							
	research collaborators from other institutions or a research sponsor). If PHI will NOT be released							
	outside of HPH, please make a statement to that effect.							
9.	In substitute will the DIII be maintained?	on on Delectronic Deth						
10.	In what form will the PHI be maintained? Paper Electronic Both							
10.	If the information is in paper format, describe the precautions you have to protect the PHI from							
	improper use and disclosure:							
11.	If data is stored electronically (PC_lapton_CD_F	OVD thumb drive portable storage device etc.)						
	If data is stored electronically (PC, laptop, CD, DVD, thumb drive, portable storage device, etc.), what safeguards are in place to prevent access to the electronic files?							
	Password protected PC CD							
	128 bit Encryption Laptop	DVD						
	Other (please specify):	Portable storage device (type)						
12.	Is access to the information restricted to only tho	se who have a "need to know" for Yes No						
	performance of their job?							
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13.	Is this electronic system used to transmit data outside of your site?	Yes	No			
14.	If information is transmitted, what safeguards are in place to prevent inadvertent access to this data during transmission?					
15.	Will you be retaining any identifiable information on potential subjects who do not meet study eligibility requirements? If yes, explain the purpose of retaining the information.	Yes 🗌	No			
16.	When do you plan to destroy the identifiers? (Identifiers must be destroyed at the earliest opportunity.) Subject Contact Enrollment Other (please specify): Describe your plan for destroying the identifiers at or before the conclusion of the study or provide a justification for long term or permanent retention of the identifiers. Specify which identifiers and information will be destroyed. If long term retention is requested, such as maintenance of a database, specify the security measures you will use.					
17.	Please explain how your recruitment meets the following criteria: 1. a. Recruitment cannot be practicably carried out without the Partial Waiver of Authorization. 2. Recruitment cannot practicably be conducted without the participants' PHI.					
By signing this statement, I am providing written assurance that only information essential to the purpose of this research will be collected. Access to the information will be limited to the greatest extent possible. Storing data on portable media devices is highly discouraged. If I do use portable media devices, I understand any identifiable data placed on portable electronic media or other devices must be encrypted. Protected health information collected under this waiver will not be re-used or disclosed to any other person or entity. You may type your name on this form and send it electronically. Your typed name on this form will constitute your signature and agreement with the aforementioned statement.						
Signat	ture of Principal Investigator Date					