

IRB Administration Office

87 East Canfield, Second Floor Office: (313) 577-1628 Detroit, Michigan 48201 http://irb.wayne.edu/index.php

HIPAA Summary Form

This form must be in the current version and typewritten

This form must be opened and saved using Adobe or software that allows for digital signature.

Instructions: Steps for Signing a PDF Form with a Digital ID

Section A: Administrative Information

1.	Principal Investigator Name	
		The Principal Investigator must sign Section E of this form.
2.	Project Title	
3.	From which of the follo	wing institutions will you obtain Protected Health Information (PHI)?
	questionnaires or surve oversight by the IRB is (Select all that apply) Detroit Medical Ce John D. Dingell VA Karmanos Cancer	nter . Medical Center
	Other (specify):	

Written HIPAA Authorization (See NOTE 1 below)	Waiver of Authorization (See NOTE 2 below)	Limited Data Set (See NOTE 3 below)
the following condition is met:	is obtained during the informed consent pro-	
 NOTE 2: A Waiver of Authorization is an armonic of the research team will use Phone of the research team will access includes accessing medical remust still obtain written HIPAA. The research team will de-ide any PHI. The PI has attached documents. 	required if one of the following conditions is a HI, but will not obtain informed consent. If PHI for research purposes prior to the information to identify persons who are eligible to a Authorization during the informed consent partify health information for all aspects of this tation by a statistician, independent of the regy the use of PHI is minimal. Note: Documer	met: med consent process. This recruit for this study. <i>Note: The process.</i> study, and will not use or disclosesearch team, that the risk of
 and justification for determining In addition: The PI must request a waiver Age is PHI for persons over 89 	of consent using the applicable IRB forms. years of age. If requesting a Waiver of Autor ver 89. If age is not collapsed into decades.	thorization, you must collapse aç
 and justification for determining In addition: The PI must request a waiver Age is PHI for persons over 85 into decades for participants of be obtained from participants NOTE 3: A Limited Data Set may be used. PHI used and/or disclosed in the participants. 	of consent using the applicable IRB forms. by years of age. If requesting a Waiver of Autorer 89. If age is not collapsed into decades, over 89.	thorization, you must collapse ag , written HIPAA Authorization mu n street (which includes city, stat
 and justification for determining In addition: The PI must request a waiver Age is PHI for persons over 80 into decades for participants of be obtained from participants NOTE 3: A Limited Data Set may be used to be performed and/or disclosed in the zip code), elements of dates, as study identification numbers). In addition: If the PI will disclose the Limited Data John D. Dingell VA Medical Center, Ka University and the outside entity must 	of consent using the applicable IRB forms. by years of age. If requesting a Waiver of Autover 89. If age is not collapsed into decades over 89. sed if the following condition is met: his study is limited to addresses greater than	thorization, you must collapse ag, written HIPAA Authorization must refer to the street (which includes city, state, characteristic or codes (i.e. links) inversity, the Detroit Medical Ceragreement between Wayne States Data Use Agreement template

Section B: Participant Recruitment and Use of PHI

5.	Will someone with a clinical relationship contact or refer potential participants to your study? NOTE: A person with a clinical relationship should first introduce a study to potential participants. This person does not have to be a member of the research key personnel. Research that is not in a clinical setting, does not involve face-to-face recruitment (i.e. advertisements), or does not involve direct contact with participants (i.e. previously collected data), may not require that a person with a clinical relationship introduces the study. a. State who will introduce the study to potential participants and their clinical relationship(s).
	b. Justify why someone with a clinical relationship will not introduce the study to potential participants.
6.	Which of the following Protected Health Information (PHI) items obtained from Wayne State University, the Detroit Medical Center, John D. Dingell VA Medical Center, Karmanos Cancer Institute are being USED for research purposes? (Select all that apply and include for the HIPAA Authorization PHI that will be USED section) NOTE: Research uses include screening for eligibility, data collection, data analysis, and follow-up contact. Name (including Initials) Address: Street Address City, State, and/or Zip Code Elements of Dates (Birth Date, Admission Date, Date of Service, Date of Death, etc.) Telephone Number Fax Number
	 □ E-Mail Address □ Social Security Number □ Medical Record Number □ Health Beneficiary Number
	Account Numbers (Credit Card, etc.) Certificate/License Numbers Vehicle Identification/Serial Numbers Device Identification/Serial Numbers
	Website URLs Internet Protocol (IP) Addresses Biometric Identifiers (Voice, Fingerprints, etc.)
	☐ Full Face Images☐ Any Other Unique Identifying Numbers, Characteristic or Code (Linked Study Identification Numbers, etc.)

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Se	ction C: Disclosure of PHI	
7.	Will PHI be DISCLOSED to sponsors, companies hired to provide study related services, or research institutions outside of Wayne State University and its affiliates (Detroit Medical Center, John D. Dingell VA Medical Center, Karmanos Cancer Institute)?	No − go directly to Q#10Yes
	NOTE: PHI is always available to federal agencies that monitor research upon request, and it is not necessary to consider them when answering this question. These agencies include the Office of Human Research Protections (OHRP), the Food and Drug Administration (FDA), the Office for Civil Rights (OCR), and the Veteran's Administration (VA) (if applicable).	
8.	List all sponsors, companies hired to provide study related services, or research institution University and its affiliates that will receive PHI. Note these entities must also be listed for the HIPAA Authorization form to be signed by must be listed where indicated following the template instruction.	participants. These entities
	a. Describe how data will be sent: NOTE: Describe actual methods and include a plan for coding and/or encryption to	maintain confidentiality.

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9.	Which of the following Protected Health Information	
	(PHI) items are being DISCLOSED to sponsors, companies hired to provide study related services,	Justify the disclosure of each selected item below.
	or research institutions outside of Wayne State	
	University and its affiliates?	
	(Select all that apply include for the HIPAA	
	Authorization PHI that will be DISCLOSED	
	section)	
	Name (including Initials)	
	Address:	
	Street Address	
	☐ City, State, Zip Code	
	Elements of Dates (Birth Date, Admission Date,	
	Date of Service, Date of Death, etc.)	
	Telephone Number	
	Fax Number	
	- Tax Nambor	
	T M-11 Address	
	E-Mail Address	
	Social Security Number	
	Medical Record Number	
	I Medical Necola Nullipel	
	Health Beneficiary Number	

Account Numbers (Credit Card, etc.)	
Certificate/License Numbers	
Vehicle Identification/Serial Numbers	
Device Identification/Serial Numbers	
Website URLs	
Website OIVEs	
☐ Internet Protocol (IP) Addresses	
Biometric Identifiers (Voice, Fingerprints, etc.)	
Diometric identifiers (voice, i ingerprints, etc.)	
Full Face Images	
Any Other Unique Identifying Numbers,	
Characteristic or Code (Linked Study	
Identification Numbers, etc.)	
1	

Section D: Waiver of HIPAA Authorization

10.	Is a Waiver of HIPAA Aut	horization being requested for the proposed study?	Yes
			☐ No – go directly to Q#16
	a. Why is a Waiver of HI	PAA Authorization being requested?	
	☐ To screen medical red	cords for eligible potential participants	
	☐ To obtain data for a re	etrospective chart review study	
	Other (specify):	-	
11.	Describe how the proposed use and/or disclosure of PHI presents no more than minimal risk to the privacy of participants:		
12.	Explain why the research could not practicably be conducted without the Waiver of Authorization:		
13.	Explain why the research could not practicably be conducted without access to, and use of PHI:		

14.	Describe the steps taken to protect identifying information (or links to identifiers) from improper use or disclosure:			
15.	Describe the plans for destroying identifying information (or links to identifiers). Specify when identifying information will be destroyed. Provide justification if identifying information is retained:			
Se	-		ure for HIPAA Submission the Principal Investigator ONLY	
Prir	ncipal Investigator's S	ignature for HIPAA Submi	ission	
	As Principal Investigator, my signature is attesting to the accuracy of this submission and requesting approval/authorization for USE and/or Disclosure of Protected Health Information as indicated for this form.			
Sigr	nature of Principal Investi	gator	Date	
Principal Investigator's Name				
Principal Investigator's Waiver of HIPAA Authorization Statement				
			estation of the following regarding the request for	
As F othe rese IRB. entit	Waiver of HIPAA Authorization. As Principal Investigator, I assure that the information I obtain as part of this research will not be reused or disclosed to any other person or entity other than those listed on this form, except as required by law, for authorized oversight of the research project, or for other research for which the use or disclosure of PHI is approved by the Wayne State University IRB. If at any time I want to reuse this information for other purposes or disclose the information to other individuals or entities, I will seek approval from the Wayne State University IRB. Yes			
=	No N/A Waiver of HIPAA Auth	orization is not requested.		

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Section F: Detroit Medical Center (DMC) Protected Health Information (PHI) Use and Disclosure

The Wayne State University IRB/Privacy Board does not review Section D of the HIPAA Summary Form. Access to Detroit Medical Center (DMC) Protected Health Information (PHI) requires DMC review. The DMC Clinical & Translational Research Office uses Section D as part of their review.

If you have questions related to Q#16-21, please contact the DMC Clinical & Translational Research Office for assistance: https://www.dmc.org/ResearchReviewProcess

16.	Are you accessing Protected Health	Yes - answer question 17.
	Information (PHI) from the DMC EMR/Medical	No − STOP, the form is complete
	Records for your research?	
17.	Are you requesting an automated extract of	Yes If Yes, answer question 18.
	DMC Protected Health Information (PHI)?	No If No, proceed to question 19.
18.	Indicate where the information is being	
	housed and transmitted:	
	18a.) A database housed within the DMC?	Yes - answer question 18b.
	Toa.) A database floused within the Divic!	No - answer question 18c
	18b.) Describe the location where the data	NO - answer question roc
	will be housed:	
	will be floused.	
	10) - " " 11	
	18c.) Transmitted to an external organization	Yes - answer question 18d.
	(eg. Wayne State University, Karmanos),	□ No
	an external database or system?	
	18d.) Describe the location where the data	
	will be transmitted:	

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19.	Are you manually extracting DMC Protected Health Information (PHI)?	Yes - answer question 20. No - STOP, the form is complete
20.	For the initial manual PHI extraction, indicate where the information will be stored :	
	20a.) A database housed within the DMC?	Yes - answer question 20b. No - answer question 20c.
	20b.) Describe the location where the data will be stored:	
	20c) An external organization (eg. Wayne State University, Karmanos), an external database or system?	Yes - answer question 20d. No
	20d.) Describe the location where the data will be stored:	
	20e.) Are you transmitting the manually extracted PHI to an external organization (eg. Wayne State University, Karmanos), an external database or system?	☐ Yes ☐ No
21.	Indicate who will be extracting the data?	□ DMC Research Team□ WSU Research Team□ DMC Information Services

	DMC Finance Other: