

This worksheet is used for subsequent submissions for any study being conducted at WSU or one of our affiliate institutions that is under the oversight of an external IRB. An external IRB is any IRB that is not the WSU IRB. Please submit this completed worksheet along with the required attachments as instructed.

Subsequent submission modifications include (for example):

•	Change in PI & Key Personnel changes.	•	Changes to the reliance agreement: (i.e transfer of the study to another outside IRB),
•	Local context changes include HIPAA Authorization and/or Waiver	•	Changes to the protocol or consent documents that affect the local context of the study.
	Changes to the injury	lang	uage in the consent form

The IRB office will notify the PI and designated personnel via email when the modification is authorized.

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

Non-eProtocol studies submit this modification request to the appropriate External IRB email box:

NCI CIRB: <u>cirb@wayne.edu</u>	WCG IRB: wirb@wayne.edu
---------------------------------	-------------------------

All others: relyirb@wayne.edu

For eProtocol SUBMISSIONS STOP

Do Not Use or Submit this form. Use the eProtocol Amendment Form

Section A: Study Details

Date Completed:
Status of Principal Investigator (check all that apply) WSU Faculty WSU Student
DMC Staff KCI Staff
Other (specify):
Title:
Email:
r this Modification Submission

2. Sponsor/Select the External IRB that is used for t	his study:	
Sponsor Contact Information		
Sponsor's Name:		Phone:
Contact Name:		Title:
If study has NIH funding, please check all that apply:	plicants 🗌 Career developme	ent applicants
Fellowship applicants N/A		
	☐ National Cancer Institute C	IRB
□ Advarra	□ Other Reviewing IRB (inser	t Name & contact below)
Institution's Name:		
IRB Point of contact (POC)Name:		
Email:	Phone #:	
Please select all that apply and o	complete the applicable	sections.
Section B: Unanticipated If Not App	Problem/Event Reporti licable, Select N/A and go t	
Unanticipated Problem (UP)/Adverse		
Protocol Violations/Deviations submitted to the Rei Include with <u>this</u> submission all documents that are pro- communications) Is this an: Initial Report Follow-Up Report External/Reviewing IRB's Determination	0	i.e., reporting form, UP
Local Context Protocol Violations/Deviations: This includes HIPAA, key personnel, and breach of cor • STOP DO NOT SUBMIT THIS MODIFICATION • Submit the UP form available on the IRB's web Problems & Event Reporting Form	N FORM	

Section C: WSU Local Key Personnel Changes

If Not Applicable, Select N/A and go to next section N/A

Study personnel: persons engaged in the collection of data or have access to data through intervention or
interaction with the participant, including the consent process, or have access to the participant's identifiable
private information. This may include collaborators, fellows, residents, research assistants, etc.

Note: Do not list/add key personnel from other sites outside of WSU and local affiliates.

- If a financial conflict of interest exists, a Financial Conflict of Interest Detailed Disclosure form must be completed then submitted to the WSU Financial Conflict of Interest Committee. The FCOI disclosure form can be found at http://research.wayne.edu/coi/index.php.
- All key personnel are required to take the CITI training program found at www.citiprogram.org.

Key Personnel Research Role and Obligations:

- (I) Complete and maintain required human participant research training and update every three years.
- (II) Follow the direction of the Principal Investigator to adhere to the IRB approved study protocol, institutional policies, and research regulations.

Does this submission include key personn	nel changes? [Yes	🔄 No
--	----------------	-----	------

Select the personnel change type:	Deletion	Addition
-----------------------------------	----------	----------

Key Personnel Additions: complete the <u>Additional Key Personnel/Change Request Form</u> and include with this submission

Key Personnel Deletion: State the name and role of individuals being deleted in the box below.

Section D: Local Context Changes & HIPAA Documents

If Not Applicable, select N/A and go to next section N/A

	Please select all that apply and attach a <mark>highlighted or <u>tracked changes</u> copy of consent/assents form(s) along with the most recent version of the IRB approved consent form.</mark>
1.	Changes to HIPAA Summary Form (including waiver) and/or HIPAA Authorization:
	 Please submit updated HIPAA Summary Form with PI signature(s) and updated consent with HIPAA Authorization revisions.
	eProtocol submissions NOTE: eProtocol Submissions complete the eProtocol HIPAA Section.
2.	Consent: Changes to research related injury language
3.	Consent: Changes to local contact info (includes for WSU PI change)
4.	Consent: Changes to study cost
5.	Consent: Changes to study title New Study Title:

6. Change in key personnel member's conflict of interest statu	s (please provide conflict of interest plan if applicable)
7. Other Changes:	
Describe proposed changes and the rationale:	
PLEASE NOTE: If this description is not cor	nnleted the submission will be returned
Section E: Ancil	
	cable, select N/A and go to next section N/A
Ancillary Reviews Do the changes require any of the following approvals? If Ye	s, must provide approval letter with this submission
Embryonic Stem Cell Research Oversight Committee (ESCRO)	No Yes (If " Yes " provide letter)
Institutional Biosafety Committee (IBC)	No Yes (If " Yes " provide letter)
Radiation Safety Committee (RSC)	No Yes (If " Yes " provide letter)
Materials Transfer Agreement (MTA)	No Yes (If " Yes " please contact
Karmanos Cancer Institute Protocol Review & Monitoring	<u>mtainfo@wayne.edu</u>)
Committee (PRMC)	
McLaren Health Care review	No Yes (If " Yes " provide letter)
Veterans Administration	No Yes (If " Yes " provide letter)
Department of Psychiatry	No Yes (If " Yes " provide letter)
Detroit Medical Center (DMC) Review	No Yes
https://www.dmc.org/for-health-professionals/clinical-translation- research-office	If " Yes " DMC Approval can be gained concurrently, but is required for IRB Administrative authorization.
Note: Research occurring at DMC must copy <u>mmontie@dmc.org</u>	
on ALL communications with WSU IRB.	

Section F: WSU PI Change

		If Not Applicable, STOP THIS FORM IS COMPLETE
Ch		nature from the new PI, current PI, and the Department Chair. able questions, 3 (c) & (d) must be completed.
1.	The Current Principal Investigato	r Name & Signature must be provided for page 1 of this form.
2.	a. Should the current PI be:	 Removed from the study. Added as key personnel in the research role of:
	b. Is the current PI available to provide an original signature on this form?	Yes – go directly to Q#4 No – answer sub-questions below (3c & 3d) and obtain a signature from the Department Chair, Dean, or Signatory Official authorizing the PI change
	c. State why the current PI is unable to provid signature):	de an original signature (include details regarding attempts to obtain a
	d. Include documentation (e.g. e-mail) from the current PI acknowledging that a PI change is appropriate, or explain why it is not possible to obtain documentation:	Documentation from the current PI is being submitted.

4. Information for the proposed new Principal Investigator	Name of new	PI:		The proposed PI's bio- sketch or CV is attached to this modification request.
	Department			
	Address		Pager	
			E-Mail	
			Telephone	
Reason for the chang	-			
What is the proposed	I PI's professior	nal and/or educational qualifications f	or being the P	l on this study?

New Drineinel Inv		
New Principal Inv	estigator's CITI Training	
Further dir eProtocol submis	ust have completed the <u>CITI training</u> program at <u>https://www.citiprogram.org/Default.asp</u> ections and a listing of the training is available at: <u>http://irb.wayne.edu/mandatory-training.php</u> The new PI must affiliate with Wayne State University for their CITI profile. sions NOTE: For eProtocol submissions, the new PI must include their WSU Access ID for their ITI profile. Note, CITI information updates in eProtocol each business morning.	r
a) Have you taken:	HIPS RCR & Basic/Refresher Course for Human Subjects?	
Yes	No - <u>STOP</u> : do not submit this modification request until are required CITI training is complete.	
b) If CITI training was t	aken under a former name (e.g. maiden),	
What is that name?		
Is the Principal Inv	estigator's CITI Training up to date?	
	WSU New Principal Investigator Attestation & Authorized Signatory oen and save form using Adobe or software that allows for electronic signature.	
	Dringing Investigator's Cignotype and Attactation	
	Principal Investigator's Signature and Attestation	
	o you, your spouse or domestic partner, and/or dependent children have a potential and/or of interest with the sponsor of this project (including all secondary sources)?	r
real financial conflic	o you, your spouse or domestic partner, and/or dependent children have a potential and/o	r
real financial conflic NO YES (<i>if yes, ple</i> In signing as the P 1. Attest to the 2. Agrees to ac approved by 3. Agrees to ab 4. Agrees to su	o you, your spouse or domestic partner, and/or dependent children have a potential and/or of interest with the sponsor of this project (including all secondary sources)? Pase include WSU Memo of Understanding/Agreement to FCOI Management Plan) rincipal Investigator, I attest/agree to: accuracy of the information provided in this submission. cept primary responsibility for the scientific and ethical conduct of the research, as	
 real financial conflic NO YES (<i>if yes, pla</i>) In signing as the P 1. Attest to the 2. Agrees to ac approved by 3. Agrees to ab 4. Agrees to su 5. Agrees to ab 	o you, your spouse or domestic partner, and/or dependent children have a potential and/or of interest with the sponsor of this project (including all secondary sources)? Pase include WSU Memo of Understanding/Agreement to FCOI Management Plan) rincipal Investigator, I attest/agree to: accuracy of the information provided in this submission. Cept primary responsibility for the scientific and ethical conduct of the research, as the IRB. de by the IRB's policies and procedures. pomit unanticipated problem/adverse event reports in a timely manner.	

Certification for Change in PI by the Dean/Chair/Authorized Signatory		
Name of the Dean/Chair/authorized signatory		Title
	Name of College/Departme	ent/Institute/Center
Is authorized signatory's	CITI training up to date?	Yes 🗌 No
	· · ·	, and/or dependent children have a potential and/or ect (including all secondary sources)?
YES (if yes, please ind	clude WSU Memo of Understand	ling/Agreement to FCOI Management Plan)
	Dean, and Institute/Center Dir	est that I am the authorized signatory for the rector for the above noted
(b) appropriate scientific an(c) the research uses proce	be provided for the research pro d ethical oversight has been and dures consistent with sound res ound enough to yield the expect	earch design.
Signature of Chair/Dean/	Authorized signatory	Date