NCIPC Determination of Applicability of Human Subjects Regulations, Request to Classify Project as Not Involving Human Subjects or Research

Project/FOA Title: Conduct an Older Adult Mobil	lity Assessment Tool Impact	Evaluation
Science Officer(s): Gwen Bergen		Telephone: 770-488-1394
Project Officer(s): Leanna Fox	Division: DUIP	Telephone: 770-488-3915
	Ethics Verification Number:1581	
Proposed Project Dates: Starting: 08/04/2014 Enc	ling: 09/08/2016	
Award Title: Conduct an Older Adult Mobility Asse	essment Tool Impact Evaluation	and Develop a Dissemination Plan
Award Institution: BATTELLE MEMORIAL INST	TTUTE	shadows surers and the same
Funding Mechanism #		
Funding Sponsor:		
Number of Participants in Study: 1,000		
☐ Intramural or 区 Extramural		
Categories of data collection that do not constitute hengaged are listed below. Please check appropriate 1. Activity is not research. Primary intent is primprovement of programs or services. Object A. Epidemic/endemic disease/injury of control needs.	public health practice: disease/inctives focused on a specific popurontrol activity; collected data d	ajury control, surveillance, ulation. irectly relate to immediate disease
B. Routine disease/injury surveillanc for a specific health condition/disea C. Program evaluation activity; data in a specific population/setting.	se in a specific population and s	etting. (Includes disease reporting)
Justification: Please attach project goals/aims, obje	ectives, design, setting and parti-	cipants, methods, and data sources.

-OR-		
☐ II. Activity is research but does NOT involve identifiable human contribute to generalizable knowledge. ☐ A. Activity is research involving collection/analysis of decorganizations or units, which are not individual personal B. Activity is research involving data and/or specimens in	data about health facilities or other	
Justification: Please attach project goals/aims, objectives, design, setti	ing and participants, methods, and data sources	3.
-OR-		
III. Activity is research involving human subjects but CDC – in and on-site contractors (but not off-site contractors or othe intervening or interacting with participants and will NOT private data or biological specimens.	have access to identifiable (including coded	1)
Justification: Please provide a summary of CDC's role and explain the data by intervening or interacting with participants or have acceded data that have been stripped of the codes that link information to "engaged" in human subjects research. Also, please attach a su setting and participants, methods, other data sources and plans	to individuals and still be considered to not be ummary of project goals/aims, objectives, design	.0
Once local IRB approval has been obtained please forward a co Human Subjects Contact - for records keeping purposes.	opy (electronic preferred) to Jahlani Akil - the	3
Attach project description in enough detail to clarify "non-human subjetthe product.	ects", "non-research" or "not-engaged" nature	of
	nstance, investigators/project officers are expecting to the maximum extent possible the privacy	eted
Comments/Rationale: Although CDC Human Subjects (IRB) review is not required in this in to adhere to ethical principles and standards by respecting and protecti	nstance, investigators/project officers are expecting to the maximum extent possible the privacy	eted
Comments/Rationale: Although CDC Human Subjects (IRB) review is not required in this in to adhere to ethical principles and standards by respecting and protectic confidentiality and autonomy of participants. All applicable State and	nstance, investigators/project officers are expecting to the maximum extent possible the privacy	eted
Comments/Rationale: Although CDC Human Subjects (IRB) review is not required in this in to adhere to ethical principles and standards by respecting and protectic confidentiality and autonomy of participants. All applicable State and Additional Comments: Required Signatures:	nstance, investigators/project officers are expecting to the maximum extent possible the privacy	eted
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Comments/Rationale: Although CDC Human Subjects (IRB) review is not required in this in to adhere to ethical principles and standards by respecting and protectic confidentiality and autonomy of participants. All applicable State and Additional Comments: Required Signatures: Digitally signed by Karen C. Angel -S DN: c=US, o=U.S. Government, ou=HHS, ou=CDC, ou=People, cn=Karen C. Angel -S, 0.9.2342.19200300.100.1.1=1000551064 Date: 2015.06.03 12:35:24-04'00' Division Official (e.g., Director or ADS)	instance, investigators/project officers are expecting to the maximum extent possible the privacy leaders of the privacy leaders of the followed.	eted
Comments/Rationale: Although CDC Human Subjects (IRB) review is not required in this in to adhere to ethical principles and standards by respecting and protectic confidentiality and autonomy of participants. All applicable State and Additional Comments: Required Signatures: Digitally signed by Karen C. Angel -S DN: c=US, o=U.S. Government, ou=HHS, ou=CDC, ou=People, cn=Karen C. Angel -S, 0.9.2342.19200300.100.1.1=1000551064 Date: 2015.06.03 12:35:24-04'00'	nstance, investigators/project officers are expecting to the maximum extent possible the privacy laws must be followed. O6/03/2015	eted

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Funding Amount: