

Grants and Contracts Office

One Gustave L. Levy Place Box 1075 New York, NY 10029-6574 Phone: 212.

Facsimile: 212.241-3294 Email: grants@mssm.edu

NIH Foreign Component Prior Approval Form

Instructions: Please submit this form to the GCO upon request. The GCO will review and submit the form to the National Institutes of Health Institute or Center (NIH I/C) for prior approval of a foreign component. Attach additional pages if more space is needed. Please review NOT-OD-19-114 for more information regarding the NIH's policy on foreign components.

Primary Institution Name:	Icahn School of Medicine at Mount Sinai (ISMMS)		
Primary Institution Address:	1 Gustave L. Levy Place, Box 107		
	ame:		
_	nust be affiliated with this institu	tion. If investigator is not affiliated with additional clarification.	
Country in which research will b	pe conducted:		
Names of Foreign Component C	Collaborating Investigators:		
Foreign Component Address: P	rovide as much detail as possible	•	
Collaborating Investigator Teler	phone Number:	_E-mail Address:	
The information requested belo include information about studi 1. AIDS Related? 2. Involves Biodefense? 3. Description of Activities to be Please describe the activities to	w is regarding the research condi- es performed in the U.S. or at and [] Yes [] No [] Yes [] No pe Conducted at the Foreign Com- be conducted at the foreign com-	ucted at the foreign component. Do not other site.	
5. Budget: [] Attached []	If appropriate, add None which includes the total costs per		
Vertebrate Animals Include the following information 6. Vertebrate Animals?	on if vertebrate animals are involv	ved at the foreign component:	

7.	For experiments involving animals, please identify the type of animal being used, how many animals, and a brief description of what will be done to them?		
	man Subjects		
	clude the following information if human subjects are involved at the foreign component:		
	Human Subjects? [] Yes [] No If Yes to Human Subjects, either state "pending approval" or include an FWA (Federal-Wide		
Э.	Assurance) # as appropriate		
	If there is no information about Approval status, then there may be a hold on approval.		
10	. Does the research at the foreign component meet the NIH definition of a Clinical Trial? []Yes []No		
	. Involves Stem Cells? [] Yes [] No		
	. Number of subjects:		
13	. Demographics: Include age-range, gender, inclusion of special populations such as pregnant women, etc.		
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14	. How will the subjects be recruited?		
	. Will informed consent be obtained? [] Yes [] No		
16	. What will the subject participation entail?		
17	. If subjects are to be interviewed, where, how long, and how many times?		
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18	. How long they will participate?		
19	. How will subject confidentiality be protected?		
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20	. If samples are to be obtained, samples of what and how often?		
21	. Other: Use this section to provide additional information		
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