

Office of Regulatory Research Compliance Material Transfer Request Form

Provide the requested information and submit this form to mta.orrc@howard.edu. Visit the ORRC website for more information.

0	RGANIZATION AND PERSONNEL INFORMATION				
1.	Howard University willthe Material defined in this form.				
	MATERIAL PROVIDER MATERIAL REC	IPIENT			
2.	Organization Name:				
3.	Contractual Contact:				
	a. E-mail Address:				
	b. Phone Number:				
4.	PI or Scientist:				
	a. E-mail Address:				
	b. Phone Number:				
	c. Department:				
	d. Mailing Address:				
	e. City, State, Zip:				
5.	Signatory Official:				
Μ	ATERIAL AND PROJECT INFORMATION				
6.	Material to be transferred:				
7.	Quantity to transfer: When will the Material be used? Start Date: End Date:				
8.	Provide the title of the project related to the Material.				
9.	Provide a brief description of the project and how the material will be used.				
5.					
10.	Is the Material available commercially or through any other source?				
	Will University space be required to complete project activities using the Material?				
	If Yes, explain:				
12.	Will the Material be used in conjunction with another material previously received under an agreement?	YESNO			
	a. If Yes, identify that material and its source. Also, forward a copy of the agreement with this form.				
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13.	Is the Material currently or expected to be associated with a Non-Disclosure Agreement?	YES NO			
14.	Was the Material created or invented at Howard University?	🗌 YES 🗌 NO			
	a. If No, identify the origin of the Material:	_			
	b. If Yes, has the Material been disclosed to the Office of Intellectual Property and Technology?	🗌 YES 🗌 NO			
15.	Is the Howard University Investigator/scientist the creator/inventor of the Material?	🗌 YES 🗌 NO			
	a. If No, identify the original creator/inventor:	-			
16.	Is the Material relevant to any previous or pending inventions disclosed to Howard University?	YES NO			
17.	Is the Material a tool, kit, or instrument that will be used in the conduct of research?	YES NO			



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KE	G	ULATORY COMPLIANCE INFORMATION				
18.	ls t	the Material dangerous to handle, store, or use?	YES NO			
	a.	If Yes, explain:				
19.	Does the project involve or contain the following? If Yes, provide the applicable information requested.					
	a.	Human participants	🗌 YES 🗌 NO	IRB #:		
	b.	Human biological samples or substances	🗌 YES 🗌 NO	IRB #:		
		1. Are the samples or substances de-identified?	YES NO			
	c.	Making progeny, unmodified derivatives, or descendant copies of the Material	YES NO	IRB #:		
	d.	The care and use of animals	YES NO	IACUC #:		
	e.	Recombinant DNA, infectious agents, toxins, or reagents	YES NO	IBC #:		
		1. Define each with the related biosafety level and IBC protocol numb	per.			
		i	Biosafety Level#:	IBC #:		
		ii	Biosafety Level#:	IBC #:		
	f.	Radioactive material YES NO	Biosafety Level#:	IBC #:		
	g.	Radioisotopes in or on humans I YES NO	Biosafety Level#:			
	h.	An antidote that is necessary for use with the Material	YES NO			
	i.	An export-controlled agent	YES NO			
	j.	Classified research	🗌 YES 🗌 NO			
	k.	Restrictions on openness of research	🗌 YES 🗌 NO			
	I.	Product testing and evaluation (i.e., testing an expression system)	🗌 YES 🗌 NO			
	m.	Off-Campus work	🗌 YES 🗌 NO			
	n.	Any other regulatory compliance concerns related to the Material	YES NO			
		1. Explain:				
CE	RT	IFICATIONS				
My signature certifies that:						
i.	i. Project activities are consistent with University objectives and the faculty involved in the project have agreed to participate.					

- ii. The obligations and commitments described are acceptable and in accordance with University and sponsor policies.
- iii. I understand that the Principal Investigator and Department Chairperson will bear responsibility for monitoring compliance with agreement terms.

Principal Investigator:	Date Signed:
University Bio-safety Officer:	Date Signed:
Senior Compliance Officer:	Date Signed:
AVP for Regulatory Research Compliance:	Date Signed: