

APPLICATION FOR PERMIT TO IMPORT INFECTIOUS BIOLOGICAL AGENTS INTO THE UNITED STATES

FORM APPROVED OMB NO. 0920-0199 EXP DATE 8/31/2024

Guidance for completing this form is available at http://www.cdc.gov/od/eaipp/importApplication/. This form may be submitted by mail, fax, or email attachment to the Centers for Disease Control and Prevention, Import Permit Program. Mailing Address: 1600 Clifton Road NE, Mailstop A-46, Atlanta, GA 30333. Fax: 404-718-2093. E-mail: lmportPermit@cdc.gov. Telephone: 404-718-2077. *Please submit completed form only once by either email, fax, or mail*

SECTION A - Person Requesting Permit in U.S. (Permittee)								
1. Permittee's Last N	Name 2	Permittee's First Nan	ne 3. Pe	rmittee's Organization	n			
4. Physical Address	(NOT a post office box)		5.	City		6. State 7. Zip Code		
8. Permittee's Telep	hone Number		9. Permittee's Email					
10. Secondary Con	tact's Name	11. Secondary Number	ry Contact's Email N	lame				
13. Institutional Bios	safety Officer's Name	14. Institutional Telephone Num	Biosafety Officer's liber	's 15. Institutional Biosafety Officer's Email Name				
CLICK HERE TO ADD ADDITIONAL ROWS (AUTHORIZED USERS OF THE PERMIT)								
SECTION B - Sender of Imported Infectious Biological Agent(s) or Vector(s)								
1. Sender's Last Na	me	2. Sende	r's First Name 3.	Sender's Organization	on			
4. Physical Address Outside of the U.S. (NOT a post office box)5. City6. State/Providence7. Country								
8. Postal Code 9. Telephone Number 10 Email								
CLICK HERE TO	O ADD ADDITION							
			C - Shipment In					
Method(s) of Ship Commercial Carri Hand-carried by in	oment er (e.g., FedEx) ndividuals listed in Sed		Estimated Number of	Shipments [Enter nu	umeric value]			
SECTION D - Description of Infectious Biological Agent(s) and Permittee's Laboratory								
1. Intended use(s) of Diagnostic Research Clinical trials Education Production				d description of the w ur work clearly & simply. In		shed with the imported ose, objectives, methods,		
Other (please describe): 3. Will the agent(s) be propagated or cultured? X Yes X No If yes, will the total culture volume exceed 10 liters at any point? X Yes X No If yes, will the agent(s) be used to inoculate animals or arthropods? X Yes X No If yes, will this be by the aerosol route? X Yes X No						ods?		
5. Scientific name of	6. Strain (if applicable)	7. Building Location	8. Suite/Room Location	9. Laboratory	10. Storage	11. Safety Level X BSL-1		

known/suspected biological agents(s) include Genus and species

X BSL-2 X BSL-3 X BSL-4 X ABSL-1 X ABSL-2 X ABSL-3 X ABSL-4 X ACL-1 X ACL-2 X ACL-3 X ACL-4 X ABSL3Ag

CLICK HERE TO ADD ADDITIONAL ROWS (Infectious Biological Agent(s))

SECTION E - Description of Material(s) Containing the Infectious Biological Agent(s) or Vector(s) to be Imported							
1. Source of material(s) being in Infected or suspected infection in Infe	ted human ted vector :e describe):	2. Description of material(s) containing biological agent(s) (Check all that apply and provide description below) a Field-collected specimen b Laboratory derived isolate/culture b Blood/blood products c Blood/blood products d Other body fluids i Provide a detailed description of the material containing the biological agent:					
		Biosafety Measures					
1. Primary Containment to be used (Check all that apply) a None (open bench) Class I Class II, Type Class III Fume Hood Negative pressure ventilated enclosure with HEPA filtration Tother (please describe):	2. Personal Protective Measures to be used (Check all that apply) a Gloves b Protective Clothing (e.g., laboratory coat) C Goggles X Face Shield Facemask N95 or N100 Respirator X Powered Air Purifying Respirator (PAPR) Immunizations Other (please describe):	 3. Personnel Training provided (Check all that apply) Risk(s) associated with the imported biological agent(s) Hazardous Material Packing/Shipping Laboratory Standard Practices Hazardous Waste Handling/Disposal Emergency Response Procedures Spill Procedures Other (please describe): 	4. Has the permittee implemented biosafety measures commensurate with the hazard posed by the infectious biological agent, infectious substance, and/or vector to be imported, and the level of risk given its intended use? a No b Yes (Plan may be required to be submitted)				
5. Anticipated disposition of Inf material containing it) when wo a Will be retained at address b Will be transferred to location Will be destroyed (please com	isted in SECTION A on listed in SECTION G	6. If Agent(s) will be destroyed, list expected method(s) of destruction Thermal: X Onsite Autoclave X Onsite Incineration Chemical (describe chemical): Irradiation (describe energy source): X Contracted hazardous waste disposal company (name of company): Other (please describe):					

SECTIO	N G – Final Des	tination(s)	of Im	port	ed Biological Age	ent(s) or Ve	ector(s)	
1. Will the permittee transfer the ir	nported materials to	locations not I	isted ir	n Sect	ion D above. X Yes (c	omplete item	s 2-25) X I	No
2. Last Name of Recipient at Dest	3. First Name			4. Destination Organization				
5. Final Destination Address (NOT	6. City			7. State	8.	8. Zip Code		
9. Telephone Number	10. Email:							
11. Intended use(s) of imported agent(s) Diagnostic Research C Clinical trials Education Production Other (please describe): 12. Provide a detailed description of the work to be accomplished with imported agent(s) (Describe your work clearly & simply. Include background, purp objectives, methods, etc.) 13. Will the agent(s) be propagated or cultured? X Yes X No If yes, will the total culture volume exceed 10 liters at any point? X Yes X No If yes, will this be by the aerosol route? X Yes X No If yes, will this be by the aerosol route? X Yes X No							de background, purpose,	
15. Scientific name of known/suspected biological agents(s) include Genus and species	16. Strain (if appli	cable)	17. Build Loca		18. Suite/Room Location	19. Laborat ory	20. Storage	21. Safety Level X BSL-1 X BSL-2 X BSL-3 X BSL-4 X ABSL-2 X ABSL-3 X ABSL-4 X ACL-1 X ACL-1 X ACL-2 X ACL-3 X ACL-4 X ABSL3Ag
22. Primary Containment to be used (Check all that apply) None (open bench) Class I Class II, Type Class III Fume Hood Negative pressure ventilated enclosure with HEPA filtration f Other (please describe):	Measures to be used (Check all that apply) Gloves Protective Clothing Goggles X Face Shield Facemask N95 or N100 Respirator X Powered Air Purifying Respirator (PAPR) Immunizations		(Chan Rist) A Rist A Rist A Rist A Rack A La A Hance A Hance A Pro B Sp	4. Personnel Training provided (Check all that apply) Risk(s) associated with the imported biological agent(s) Hazardous Material acking/Shipping Laboratory Standard Practices Hazardous Waste andling/Disposal Emergency Response Procedures Spill Procedures Other (please describe):		25. Has the permittee implemented biosafety measures commensurate with the hazard posed by the infectious biological agent, infectious substance, and/or vector to be imported, and the level of risk given its intended use? a No Pes (Plan may be required to be submitted)		
+ CLICK HERE TO ADD ADDITIONAL ROWS (Final Destinations of Imported Biological Agent(s) or Vector(s))								
I hereby certify that all individuals listed in this application have the appropriate qualifications, experience and training to safely handle the agents being imported and that the information submitted in this application is complete and accurate to the best of my knowledge and belief. I agree to comply with all conditions, restrictions and precautions that may be specified in any permit that may be issued. Additionally, I agree to comply with all applicable regulations and guidelines that govern this transfer. I understand that failure to comply with the importation requirements may subject me to criminal penalties pursuant to 42 U.S.C. 271. I understand that any false statement made in this application may subject me to criminal penalties pursuant to 18 U.S.C. 1001.								
understand that any haise statement made in this a					of Permittee			
1. Permittee's Signature (REQUIRED) 2. Permittee's Printed Name (<i>Print name</i>) 3. Date Signed (<i>mm/dd/yyyy</i>)								

Public recording burden of this collection of information is estimated to average 20 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC Reports Clearance Officer; 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN: PRA (0920-0199)