

INCIDENT FORM TO REPORT POTENTIAL THEFT, LOSS, RELEASE, OR OCCUPATIONAL EXPOSURE (APHIS/CDC FORM 3)

FORM APPROVED OMB NO.0579-0213 OMB NO. 0920-0576 EXP DATE 12/31/2018

INSTRUCTIONS

Detailed instructions are available at http://www.selectagents.gov/form3.html. Answer all items completely and type or print in ink. This report must be signed and submitted to either APHIS or CDC:

Animal and Plant Health Inspection Service Agriculture Select Agent Services 4700 River Road Unit 2, Mailstop 22, Cubicle 1A07 Riverdale, MD 20737 FAX: (301) 734-3652

E-mail: AgSAS@aphis.usda.gov

Centers for Disease Control and Prevention Division of Select Agents and Toxins 1600 Clifton Road NE, Mailstop A-46 Atlanta, GA 30329

FAX: (404) 471-8375 E-mail: form3@cdc.gov

Accession Number:	ı
(For Program Use ONLY)	

Submit completed form only once by either e-mail, fax, or mail

	SECTION 1 – TO BE COMPLETED BY ALL ENTITIES							
1. Da	te of Incident:	2. Date of Immediate Notification:		3. Type	3. Type of Immediate Notification:			
					☐ E-mail ☐ Fax ☐ Telephone			
Name of Entity (entities registered with CDC or APHIS) or				5. Entity Registration Number (For select agent registered entities			gent registered entities	
Name of Hospital or Laboratory (non-registered entities):				only):				
6. Physical Address: 7. City:						8. State:	9 7ir	o Code:
U. Friysical Address.					o. State.	7. 21	o couc.	
10. Responsible Official (registered) or Name of Laboratory Supervisor (non-registered):								
11. T	elephone #:	12. Fax #:		13. E-mail address:				
14a: Type of Incident:			15. Did the release result in a potential exposure?					
☐ Theft ☐ Loss ☐ Release			☐ No ☐ Yes ☐ N/A (If Yes, explain in Blocks 28 or 30)					
☐ Unintended Animal Infection ☐ Unintended Plant Agent Release ☐ Did			Did the release result in a laboratory-acquired infection?					
Other			☐ No ☐ Yes ☐ N/A (If Yes, explain in Blocks 28 or 30)					
14b: Transfer:			If you had madical symptillance had initiated?					
Transfer incident (complete Sections 1 and 2 and Appendix B)			If yes, has medical surveillance been initiated?					
			☐ No ☐ Yes ☐ N/A (If Yes, explain in Blocks 28 or 30)					
16. Ti	me incident occurred:	17. Location of incident (buil	ding and roon		18. Locati centrifuge		n room	(e.g., freezer, incubator,
	osafety level: SL2 BSL3 BSL4	20. Date of last inventory (for reporting los only):		SS 2	21. Name of Principal Investigator:			
	BSL2 ABSL3 ABSL4 PQ Agent BSL3Ag							
		SECTION 2 – TO BE COI	MPLETED I	BY ALL	ENTIT	IES		
22. Name of Select Agent or Toxin					racterization of Age j., strain, ATCC #)	ent	24. Quantity / Amount	
Α								
В								
С								

root cause can be identified. State specifically what personal protective equiprincident involves a non-human primate, please state species. For discovery c	d. Whenever possible, conduct a risk assessment of the event and determine if the ment was worn and what, if any, medical surveillance was provided or planned. If if select agents and toxins in unregistered locations, include your entity's plan of I disposition of the discovered agents, inventory reconciliation and assurance that the , or release.
Block 25. Continued: (Use Appendix A for continuation, if necessary)	BY ALL ENTITIES ONLY FOR RELEASE
OF SELECT AGENTS AND TOX	INS OR OCCUPATIONAL EXPOSURE
26. An internal review of laboratory procedures and policies has been initiated toxins at this entity.	to lessen the likelihood of recurrences of theft, loss or release of select agents and
☐ No ☐ Yes If yes, please provide additional de	tails.
27. What were the hazards posed to humans by the extent of the release or o	ccupational exposure?
28. What is the estimated extent of the release or exposure in relation to the p	roximity of susceptible humans, animals, and plants?
29. Provide a brief summary of how the laboratory and work surfaces were de	contaminated after the release
27. Flovide a bilet sufficially of flow the laboratory and work surfaces were de	Contaminated after the release.
30. In select agents and toxins posing a risk to humans, please state how mar surveillance provided (do not provide names or confidential information).	ny laboratorians were potentially exposed and provide a brief summary of the medical
Certification: I hereby certify that the information contained on this form is true an false statement on any part of this form, or its attachments, I may be subject to conselect agent regulations may result in civil or criminal penalties, including imprison	riminal fines and/or imprisonment. I further understand that violations of the
Signature of Respondent:	Title:
Typed or printed name of Respondent:	Date:

APPENDIX A ADDITIONAL SHEET FOR CONTINUATION OF INFORMATION			
Continue Form 3 comments here. State which block from the Form 3 the continuation is from. (Example: The following statement is a continuation of block 25:):			
	Continue on next page		

APPENDIX A ADDITIONAL SHEET FOR CONTINUATION OF INFORMATION	

APPENDIX B IF THE INCIDENT OCCURRED DURING TRANSFER, COMPLETE SECTIONS 1 AND 2 OF FORM 3 AND PROVIDE THE FOLLOWING INFORMATION (INCLUDE A COPY OF THE RELEVANT APHIS/CDC FORM 2)			
Transfer authorization number from APHIS/CDC Form 2:	2. Date Shipped:		
3. Name of Carrier:	4. Airway bill number, bill of lading number, tracking number:		
5. Package Description (size, shape, description of packaging in	cluding number and type of inner packages; attach additional sheets as necessary):		
Package with select agents and toxins received by requestor:	7. Package with select agents and toxins appears to have been opened:		
☐ No ☐ Yes If yes, date of receipt:	☐ No ☐ Yes If yes, include explanation in box 5 above.		
8. Sender was contacted regarding incident:	9. Carrier/courier was contacted regarding incident: 9. Carrier/courier was contacted regarding incident:		
□ No □ Yes	□ No □ Yes		
	orm is true and correct to the best of my knowledge. I understand that if I knowingly provide a e subject to criminal fines and/or imprisonment. I further understand that violations of the uding imprisonment. 7 CFR 331, 9 CFR 121, 42 CFR 73.		
Signature of Respondent:	Title:		
Typed or printed name of Respondent:	Date:		

Public reporting burden: Public reporting burden of providing this information is estimated to average 1 hour 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/ATSDR Reports Clearance Officer; 1600 Clifton Road NE, MS D74, Atlanta, Georgia 30329; ATTN: PRA (0920-0576).