

Virus Deposit Information Form

Section 1: Type of deposit

Type of deposit	□ Catalogue: by ticking this box you declare that you understand that cultures of the deposited virus will be listed online, marketed, and distributed by NCPV for research use only under UKHSA Terms and Conditions of Supply (www.culturecollections.org.uk/orderinginfo/terms)
	□ Safe
	Patent

Section 2: Depositor information

Title and name	
Institution	
Address	
Email address	
Telephone	

Section 3: Virus identity and characteristics

Virus name in full		Titre (if known)	
Strain		Serotype / Subtype	
Virus family		Nucleic acid type and sense	
Sample type	 Supernatant Viable infected cells Cell lysate Purified virus Other (give details) 	Ampoule label	
Volume per ampoule		Number of ampoules	
Characteristics of note			
Recommended identification method	 See details (give details) See reference (give details) See attached 		



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Section 4: Safety information

ACDP hazard group	See the Advisory Committee on Dangerous Pathogens (ACDP) Approved List of Biological Agents at <u>https://www.hse.gov.uk/pubns/misc208.pdf</u>		
SAPO hazard group	See the Guidance for licence holders on the containment and control of specified animal pathogens at https://www.hse.gov.uk/pubns/priced/hsg280.pdf		
Is the virus listed in Schedule 5 Pathogens and Toxins to the UK Anti- terrorism, Crime and Security Act 2001?	□ No □ Yes		
Please supply your BSDS/COSHH form	□ Attached □ To follow		
What disinfectant is used when working with this virus?	Please include dilution and contact time (plus published literature/guidance if known)		
Brief description of known pathogenicity of the virus	e.g. Ability of this virus to survive, establish and disseminate in the environment. Include pathogenicity to humans and other natural hosts		
Could other pathogens be present in the culture?	□ No □ Yes (give details)		
Is the virus genetically modified?	 □ No □ Yes Method of genetic modification, identity and source of gene introduced, risk of conferring pathogenic traits to host and related organisms 		

Section 5: Recommended culture conditions – for catalogue items only

Cells & infection conditions	e.g. cell type, confluency at infection, MOI		
Culture medium, growth requirements	e.g. HEPES/CO ₂ , serum, incubation temperature		
Incubation time		Harvest method	
Typical CPE observed			
Long-term storage conditions			



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Section 6: Origin and history since isolation – for catalogue items only

Where did you obtain the virus from?	 Isolated in my laboratory From another laboratory (give details) Unknown 	
Original isolation information	e.g. Date isolated, cell type, incubation time, CPE observations, investigator, and institution	
Original clinical sample information	e.g. Date collected, location, sample type, patient gender, age, clinical details, travel history	
How is this sample related to the original isolate?	Include passage history, if known	
Reference for citation of the virus		
Important note: We must be able to confirm whether this organism falls within or out of scope of the Nagoya Protocol, e.g. organisms will be out of scope if isolated prior to 2014 and/or depending on country of origin.		

Section 7: Declaration

I confirm that the details given here are full and true to the best of my knowledge.

Any additional conditions of supply?	□ No □ Yes (please attach)		
Signature		Date	
Please complete and return this form before sending the virus. Please contact us if you require any assistance.			
CultureCollections.NCPV@ukhsa.gov.uk			

For Culture Collections use only

	Catalogue	ACDP Hazard Group	
Type of Deposit	🗆 Safe	SAPO? Schedule 5?	
	Patent	ACDP Containment Level	
CBA-1 notification:	 Not required Required Sent: Received: 	GMSC notification:	 Not required Required Sent: Received:
Possibility of restricted		Import licence needed to	
plant pathogen?		receive from depositor?	
Approval signature:		Approval date:	
Accession number(s):		Batch number(s):	