Australian Government
Department of Agriculture, Fisheries and Forestry
Australian Quarantine and Inspection Service

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File Ref: 2012/08444

Quarantine Act 1908 Section 13(2AA)

Permit to Import Quarantine Material

Permit: IP12009939 **Valid From:** 28 Jun 2012 **Valid To:** 28 Jun 2014

Importer	Exporter
Dr Alamgir Khan Australian Proteome Analysis Facility (APAF) Level 4 Building F7B Research Park Drive Macquarie University NSW 2109 Attn: Dr Alamgir Khan	Various Suppliers Exporters Various Addresses In All countries

You are authorised to import the following material under the listed conditions

Note: This permit covers AQIS quarantine requirement only.

All imports may be subject to quarantine inspection on arrival to determine compliance with the listed permit conditions and freedom from contamination. Imports not in compliance or not appropriately identified or packaged and labelled in accordance with the import conditions they represent may be subject to seizure, treatment, re-export or destruction at the importer's expense.

Additionally, all foods imported into Australia must comply with the provisions of the *Imported Food Control Act 1992*, and may be inspected and/or analysed against the requirements of the Australia New Zealand Food Standards Code.

All imports containing or derived from Genetically Modified material must comply with the *Gene Technology Act 2000*.

It is the importer's responsibility to identify, and to ensure it has complied with, all requirements of any other regulatory organisations and advisory bodies prior to and after importation including The Australian Customs Service, The Department of Health and Ageing, Therapeutic Goods Administration, Australian Pesticides and Veterinary Medicines Authority, Department of the Environment, Water, Heritage and the Arts, Food Standards Australia New Zealand and any state agencies such as Departments of Agriculture and Health and Environmental Protection authorities. Importers should note that this list is not exhaustive.

This permit is granted for the purposes of the *Quarantine Act 1908* and *Quarantine Proclamation 1998* of the Commonwealth of Australia. The laws of Australian States and Territories may also impose restrictions on the import of animals, plants and other goods into those States and Territories. This import permit does not prevent the application of those State and Territory laws. The importer should seek its own advice on any restrictions that may apply in any State or Territory into which it is proposed to import the animals, plants or other goods to which this permit relates.

Import conditions are subject to change at the discretion of the Director of Quarantine. This permit may be revoked without notice.

Notification of the import must be provided to AQIS for all imported goods other than goods imported as accompanied baggage or goods imported via the mail and not prescribed under the *Customs Act 1901*. Notification must be consistent with *Quarantine Regulations 2000* (examples include a Quarantine Entry or a Quarantine declaration).

Commodity Name	Condition Number(s)	Country	End Use
Animal fluids and tissues (excluding reproductive material) - sourced from avians only.	PC0992 AND PC0701 AND PC1648 AND PC6485	All countries	In-vitro use or in-vivo use in laboratory organisms only

This permit is granted subject to the condition that fees determined under Section 86E are paid

Stamp:

Delegate of Director of Quarantine

Printed Name Nathan Jamieson **Date** 28 Jun 2012

Commodity Name	Condition Number(s)	Country	End Use
Animal fluids & tissues (excluding reproductive material) - from ovines, caprines, bovines, cervines	PC0992 AND PC0701 AND PC1649 AND PC6485	All countries	In-vitro use or in-vivo use in laboratory organisms only
Animal fluids & tissues (excluding reproductive material) - sourced from equines only.	PC0992 AND PC0701 AND PC1650 AND PC6485	All countries	In-vitro use or in-vivo use in laboratory organisms only
Animal fluids & tissues (excluding reproductive material) - sourced from porcines only.	PC0992 AND PC0701 AND PC1651 AND PC6485	All countries	In-vitro use or in-vivo use in laboratory organisms only
Animal fluids and tissues (excluding reproductive material) - listed in PCT1101	PCT1101 AND PC0992	All countries	In-vitro use or in-vivo use in laboratory organisms only
Animal fluids and tissues (excluding reproductive material) - sourced from captive primates only.	PC0992 AND PC0701 AND PC0714 AND PC6485	All countries	In-vitro use or in-vivo use in laboratory organisms only

Condition	Condition Text
PC0701	PACKAGING REQUIREMENTS 1. The products must be imported in quantities of no greater than 20 mL or 20 g for each individually packaged unit.
PC0714	1. A valid copy of this Import Permit (or a method of identifying the Import Permit such as the Import Permit number) and all required documentation must accompany each consignment. Alternatively, necessary documentation will need to be presented to AQIS at the time of clearance. In order to facilitate clearance, airfreight or mail shipments should have all documentation securely attached to the outside of the package, and clearly marked "Attention Quarantine". Documentation may include Import Permit (or Import Permit number), manufacturer's declaration and invoice. The importer must meet all costs associated with the importation of this product. DECLARATION REQUIREMENT 2. Each consignment must be accompanied by a manufacturer's declaration, stating: The fluids (including urine and faeces) and tissues were obtained from primate held captive in a laboratory or zoological facility only. These primates were not known to infected with a disease agent. The manufacturer's declaration must be: .from The Manufacturer . where the manufacturer is not specified above, the declaration must be issued by the individual manufacturing site or by the manufacturer's head office within the country of export. . on manufacturer's letterhead including company address and country.

Condition	Condition Text
	. written in English. . signed by a designated representative whose name, position and title also appear. . identify the date of issue. . issued and dated within the last 6 months (unless otherwise specified in this import permit). . free from erasures and non certified alterations (all erasures and alterations must be endorsed by the issuer of the document. The only acceptable endorsement is a company stamp or seal and the signature of the company officer responsible for signing the declaration applied adjacent to the alteration). . contain the correct statement/s as required by the import conditions (all prescribed information on the certification must be legible and appear above the signature). . specific to the product(s) listed on this permit. . have a unique identifiable link to the consignment such as one of the following: container number, bill number, commercial invoice number, preferential tariff certificate number, health certificate number, packing list number or letter of credit number, batch/serial number or date of manufacture. All documentation must meet the requirements of the Minimum Documentary Requirements Policy. For full details of the AQIS minimum documentary requirements, please refer to http://www.daff.gov.au/aqis/import/general-info/documentary-requirements.
PC0992	1. A valid copy of this Import Permit (or a method of identifying the Import Permit such as the Import Permit number) and all required documentation must accompany each consignment. Alternatively, necessary documentation will need to be presented to AQIS at the time of clearance. In order to facilitate clearance, airfreight or mail shipments should have all documentation securely attached to the outside of the package, and clearly marked "Attention Quarantine". Documentation may include Import Permit (or Import Permit number), and invoice. DOCUMENTATION REQUIREMENTS 2. Each consignment must be clearly identified and linked to the relevant item(s) on the Import Permit. Identifying documentation must be available to the quarantine officer at the time of clearance. This documentation may include: a) an accompanying invoice or airway bill; or b) the physical labelling of the goods; or c) an overseas supplier's declaration describing the goods. 3. If the product description on the Import Permit varies from the identifying documentation provided for clearance, the importer is responsible for providing evidence to the quarantine officer that the Import Permit covers the products in the consignment. 4. Providing all documentation is in order at the time of clearance, the consignment can be released from quarantine. POST ENTRY / END USE CONDITIONS 5. This Import Permit allows for the importation of goods for in vitro laboratory studies (or in vivo

Condition	Condition Text
	use in laboratory organisms only), unless approved by AQIS for specific in vivo use in non-laboratory organisms. 6. This Import Permit does not permit the use of the samples for microbiological cultures or viral isolation, without prior written approval from AQIS. 7. Laboratory organisms include those defined in the following list and must be contained under laboratory or animal house conditions (or equivalent): guinea pigs, hamsters, mice, rabbits, rats, rodents or micro-organisms. Work in all other animals and plants is not permitted. 8. For in vivo use in non-laboratory organisms (eg. chickens, sheep, cattle, etc.) or plants a separate application for in vivo use must be lodged with, and approved by AQIS. This also applies if the product is to be used in veterinary vaccine or veterinary therapeutic manufacture. 9. This Import Permit does not permit the direct or indirect exposure of the imported materials or derivatives to non-laboratory organisms or plants. 10. It is the importer's responsibility to ensure that the goods are labelled "In vitro use or in vivo use in laboratory organisms only" or equivalent on the smallest packaged unit prior to distribution. 11. It is the importer's responsibility to ensure compliance with all international (eg IATA) and domestic requirements concerning the safe handling, transport and labelling of biological material. 12. It is the end user's responsibility to ensure that all laboratory products are used in accordance with the current AS/NZS 2243 Safety in Laboratory standards and Office of Gene Technology Regulator (OGTR) requirements.
PC1648	AVIAN SOURCING/POST ENTRY REQUIREMENTS 1. The product must be sourced from one of the following countries: a) Argentina, Australia, British Virgin Islands, Canada, Chile, Cyprus, Fiji, Finland, Greece, Greenland, Holland, Iceland, Republic of Ireland, Israel, Luxembourg, Malta, New Caledonia, New Zealand, Norway, Papua New Guinea, Singapore, Spain, United Kingdom/Guensey, Isle of Man, Jersey, United States of America, Vanuatu, Falkland Islands. OR b) If the product is not sourced from one of the above countries it must be subjected to gamma irradiation at 50 kGy (5 Mrad) prior to release from Quarantine. Irradiation at 50 kGy at an AQIS approved facility is mandatory even if the product has been irradiated prior to import into Australia.

Condition	Condition Text
PC1649	**Ovine, caprine, bovine, cervine and camelid.** **Sourcing and post entry requirements** 1. The product must be sourced from one of the following countries: a) Australia, Austria, Belgium, British Honduras, British Virgin Islands, Canada, Chile, Cook Islands, Denmark, Falklands Island, Fiji, France, Finland, French Polynesia, Germany, Greenland, Republic of Ireland, Italy, Luxembourg, Iceland, Malta, Mauritius, Mexico, The Netherlands, New Caledonia, New Zealand, Norway, Papua New Guinea, Portugal, Singapore, Spain, Sweden, Switzerland, United Kingdom (excluding products sourced after 1 July 2007 and before 19 February 2008), United States of America, Vanuatu. **or** b) If the product is not sourced from one of the above countries it must be subjected to gamma irradiation at 50 kGy (5 Mrad) prior to release from Quarantine. Irradiation at 50 kGy at an AQIS approved facility is mandatory even if the product has been irradiated prior to import into Australia.
PC1650	EQUINE SOURCING/POST ENTRY REQUIREMENTS 1. The product must be sourced from one of the following countries: a) Argentina, Australia, Austria, Belgium, British Honduras, British Virgin Islands, Canada, Chile, Cook Islands, Cyprus, Denmark, Fiji, Finland, France, French Polynesia, Germany, Greece, Greenland, The Netherlands, Iceland, Republic of Ireland, Israel, Italy, Japan, Luxembourg, Malaysia, Malta, Mauritius, Mexico, New Caledonia, New Zealand, Norway, Papua New Guinea, Portugal, Singapore, Spain, Sweden, Switzerland, United Kingdom, United States of America, Vanuatu, Falkland Islands. OR b) If the product is not sourced from one of the above countries it must be subjected to gamma irradiation at 50 kGy (5 Mrad) prior to release from Quarantine. Irradiation at 50 kGy at an AQIS approved facility is mandatory even if the product has been irradiated prior to import into Australia.
PC1651	PORCINE SOURCING/POST ENTRY REQUIREMENTS

Condition	Condition Text
	1. The product must be sourced from one of the following countries: a) Australia, Austria, Belgium, British Honduras, British Virgin Islands, Canada, Chile, Cook Islands, Cyprus, Denmark, Fiji, Finland, French Polynesia, Greenland, The Netherlands, Iceland, Republic of Ireland, Malta, Mauritius, New Caledonia, New Zealand, Norway, Papua New Guinea, Singapore, Spain, Sweden, United Kingdom (excluding products sourced after 1 July 2007 and before 19 February 2008), United States of America, Vanuatu, Falkland Islands. OR b) If the product is not sourced from one of the above countries it must be subjected to gamma irradiation at 50 kGy (5 Mrad) prior to release from Quarantine. Irradiation at 50 kGy at an AQIS approved facility is mandatory even if the product has been irradiated prior to import into Australia.
PC6485	ANTISERA This permit also allows for the import of anti-sera from these species. If importing anti-sera the following additional condition applies: 1) The anti-sera may only be raised against synthetic material or against antigens derived from multicellular organisms. Anti-sera raised against microorganisms (including viruses) and prions are not permitted under these conditions.

End of Condition Text

Delegate of Director of Quarantine
Printed Name Nathan Jamieson **Date** 28 Jun 2012

PCT1101

Animal fluids, tissues (excluding reproductive material) and anti-sera as listed below.

1. Animal fluids and tissues sourced from all species (excluding salmonid fish, non-human primates, humans, avians, ovines, caprines, bovines, cervines, equines, porcines, camelids or giraffids); if imported in quantities of no greater than **20ml or 20g** per smallest packaged unit

2. Anti-sera derived from all species (excluding salmonid fish, non-human primates, humans, avians, ovines, caprines, bovines, cervines, equines, porcines, camelids or giraffids); if imported in quantities of no greater than **20ml or 20g** per smallest packaged unit. The anti-sera must only be raised against synthetic material or against antigens derived from multicellular organisms.

3. Urine sourced from all species (excluding salmonid fish, non-human primates, humans, avians, ovines, caprines, bovines, cervines, equines, porcines, camelids or giraffids); if imported in quantities of no greater than **500ml or 500g** per smallest packaged unit.

4. Animal fluids and tissues sourced from all species and dried onto filter paper, dip sticks or swabs.

Notes:

- Fluids and tissues include all fluids produced by and all tissues derived from the animals specified above e.g. blood (and blood products including sera), milk, urine, faeces, mucus etc; with the exception of reproductive material.
- Reproductive material is specifically excluded from this case.
- Anti-sera raised against microorganisms (including viruses) and prions is not permitted under this case.
- For fixed tissues please refer to ICON