Australian Government
Department of Agriculture

Quarantine Act 1908 Section 13(2AA)

Phone: 02 6272 4578
Fax: 02 6272 5991
File Ref:

Permit to Import Quarantine Material

Permit: IP15009613 **Valid From:** 28 Jul 2015 **Valid To:** 28 Jul 2017

Importer	Exporter
Dr David Sayer Conexio Genomics Pty Ltd 20 Collie St Fremantle WA 6160	Various Suppliers Exporters Various Addresses In All countries

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

Note: This permit covers the Department of Agriculture 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 and Border Protection Service, The Department of Health and Ageing, Therapeutic Goods Administration, Australian Pesticides and Veterinary Medicines Authority, Department of Sustainability, Environment, Water, Population and Communities, 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.

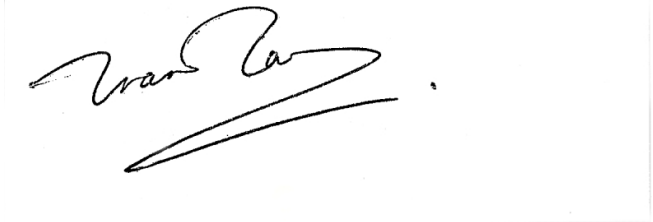
This permit is granted for the purposes of the *Quarantine Act 1908* and *Quarantine Proclamation 1998* of the Commonwealth of Australia. The laws of Australian States and Territories may also impose restrictions on the import of animals, plants and other goods into those States and Territories. This 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 the Department of Agriculture 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).

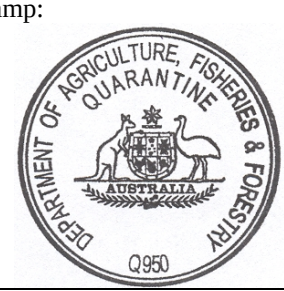
Commodity Name	Condition Number(s)	Country	End Use
Genetic material and vectors - low risk	PC0017 AND PC6797 AND PC5887	All countries	In-vitro use or in-vivo use in laboratory organisms only
Microorganisms, including viruses (as listed in PC6800)	PC0017 AND PC6800 AND PC6760 AND PC0691	All countries	In-vitro use or in-vivo use in laboratory organisms only
Genetic material - purified & derived from microorganisms & viruses (excluding high risk species)	PC0017 AND PC6855 AND PC6805 AND PC6856 AND PC0992	All countries	In-vitro use or in-vivo use in laboratory organisms only
Human fluids & tissues - free from listed diseases	PC0017 AND PC4206	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



Delegate of Director of Quarantine
Printed Name Tran Tang **Date** 28 Jul 2015

Stamp:



Commodity Name	Condition Number(s)	Country	End Use
Antibodies - Purified & raised against antigens from multicellular organisms or synthetic material	PC0017 AND PC0992 AND PC0701	All countries	In-vitro use or in-vivo use in laboratory organisms only
Human Diagnostic Tests and Human Diagnostic kits (excluding those testing for high risk human pathogens and diseases)	PC0017 AND PC6792 AND PC6369 AND PC6806	All countries	In-vitro
Diagnostic and Research Only Kits - not testing for microorganisms, viruses or prions - use in lab	PC0017 AND PC6282 AND PC0766	All countries	In-vitro
Purified / refined laboratory reagents - low risk laboratory material	PC0017 AND PC6799 AND PC0992 AND PC0701	All countries	In-vitro use or in-vivo use in laboratory organisms only

Condition	Condition Text
PC0017	**Biological Imports Program (BIP) - Administrative conditions**

1. This import permit (or number) and all required documentation must accompany each consignment and must be valid at the time the cargo is landed.

2. In order to facilitate clearance of mail shipments, the import permit (or number) and all documentation should be securely attached to the outside of the package and marked 'Attention Quarantine'.

3. The importer must meet all costs associated with the import of this product.

4. The importer (or agent) must lodge a quarantine entry for each consignment.

5. Documents must be provided with each consignment which:
a) identify the consignment e.g. entry number: and
b) identify all goods being imported as part of this consignment e.g. invoice or waybill or importers manifest; and
c) describe the goods being imported (where not clear) Example 1: Product XRab = Purified protein derived from rabbits. Example 2: Product AX = Synthetic antibiotic. Example 3: Comte = Cheese.
Note: It is the importer's responsibility to provide any additional information which is requested in order to demonstrate that the import permit covers all goods being imported.

6. Consignments that do not meet the import conditions will remain under the Department's control pending export or destruction at the importers expense.

7. For further information please contact:

Regional - Clearance assistance: http://www.daff.gov.au/biosecurity/about/contact/regional

Canberra - Biological Import Program - Administrative assistance:

Condition	Condition Text

bioadmin@agriculture.gov.au

Canberra - Biological Import Program – Technical assistance: biologicals@agriculture.gov.au

PC0691 **Packaging Requirements**

1. Cultures must be pure cultures (unless otherwise specified by this Import Permit) and labelled with the scientific name of the organism as it appears on this Import Permit.

Post Entry Requirements

2. This Import Permit allows for the importation of goods for in vitro laboratory studies (or in vivo use in laboratory organisms only), unless approved by the Department of Agriculture for specific in vivo use in non-laboratory organisms.

3. Laboratory organisms are those defined in the following list and must be contained under laboratory or animal house conditions: guinea pigs, hamsters, mice, rabbits, rats or micro-organisms. Work in all other animals and plants is not permitted.

4. For in vivo use in non-laboratory organisms (e.g. chickens, sheep, cattle, etc.) or plants a separate application for in vivo use must be lodged with, and approved by the Department of Agriculture. This also applies if the product is to be used in veterinary vaccine or veterinary therapeutic manufacture.

5. 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. This includes handling and disposal procedures.

6. It is the importer's responsibility to ensure compliance with all international (e.g. IATA) and domestic requirements concerning the safe handling, transport and labelling of biological material.

7. It is the end user's responsibility to ensure that all laboratory products are used in accordance with the Office of the Gene Technology Regulator (OGTR) requirements.

PC0701 PACKAGING REQUIREMENTS

The products must be imported in quantities of no greater than 20 mL or 20 g for each individually packaged unit.

PC0766 **Laboratory material for in vitro use only**

Post entry / end use conditions

1. This Import Permit allows for the importation of goods for in vitro laboratory studies only.

2. This Import Permit does not permit the use of the samples for microbiological cultures or viral isolation.

3. This Import Permit does not permit the direct or indirect exposure of the imported materials or

Condition	Condition Text

derivatives to animals (including laboratory animals) or plants.

4. For in vivo use in animals (including laboratory animals) or plants a separate application for in vivo use must be lodged with, and approved, by the Department of Agriculture.

5. It is the importer's responsibility to ensure that the goods are labelled "In vitro use only" on the smallest packaged unit prior to distribution.

6. 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.

7. 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.

PC0992 **POST ENTRY / END USE CONDITIONS**

1. This Import Permit allows for the importation of goods for in vitro laboratory studies (or in vivo use in laboratory organisms) only.

2. Laboratory organisms are those defined in the following list and must be contained under laboratory or animal house conditions: guinea pigs, hamsters, mice, rats, rabbits or micro-organisms. Work in all other animals and plants is not permitted.

3. For all uses in non-laboratory organisms (e.g. chickens, sheep, cattle, etc.) or plants a separate application for in vivo use must be lodged with, and approved by the Department of Agriculture. This also applies if the product is to be used in veterinary vaccine or veterinary therapeutic manufacture.

4. This Import Permit does not permit the direct or indirect exposure of the imported materials or derivatives to non-laboratory organisms or plants.

5. This Import Permit does not permit the use of the samples for microbiological cultures or viral isolation.

6. It is the importer's responsibility to ensure that the goods are labelled "In vitro use or in vivo use in laboratory organisms only" on the smallest packaged unit prior to distribution.

7. It is the importer's responsibility to ensure compliance with all international (e.g. IATA) and domestic requirements concerning the safe handling, transport and labelling of biological material.

8. 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.

Condition	Condition Text
PC4206	**Human fluids & tissues**

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 Department of Agriculture 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.

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) a manufacturers 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.

Manufacturer's declaration

4. The consignment must be accompanied by a declaration from the manufacturer stating that, to the best of my knowledge, the specimens were only taken from persons with no signs or symptoms of the following diseases:

- Cholera
- Highly pathogenic avian influenza in humans (HPAIH)
- Human swine influenza with pandemic potential
- Plague
- Rabies
- Severe acute respiratory syndrome (SARS)
- Small pox
- Viral haemorrhagic fever in humans
- Yellow fever

Condition	Condition Text

· Any disease that is exotic to Australia

5. If the consignment is not accompanied by the above declaration or the specimens are known to be infected, or are potentially infected with a human quarantinable or exotic disease, the material must be subjected to gamma irradiation at 50 kGy (5 Mrad), prior to release from Quarantine. Irradiation at 50 kGy at a Department of Agriculture approved facility is mandatory even if the product has been irradiated prior to import into Australia.

6. Providing all documentation is in order at the time of clearance, the consignment can be released from quarantine.

The manufacturer's 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.
. 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 Department of Agriculture minimum documentary requirements, please refer to http://www.daff.gov.au/biosecurity/import/general-info/documentary-requirements.

Post entry conditions

7. This Import Permit allows for the importation of goods for in vitro laboratory studies (or in vivo use in laboratory organisms only), unless approved by Department of Agriculture for specific in vivo use in non-laboratory organisms.

8. 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.

9. 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 Department of Agriculture. This also applies if the product is to be used in veterinary vaccine or veterinary therapeutic manufacture.

10. This Import Permit does not permit the direct or indirect exposure of the imported materials or

Condition	Condition Text

derivatives to non-laboratory organisms or plants.

11. 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.

12. 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.

13. 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 the Gene Technology Regulator (OGTR) requirements.

PC5887 **POST ENTRY / END USE CONDITIONS**

1. This Import Permit allows for the importation of goods for in vitro laboratory studies (or in vivo use in laboratory organisms) only.

2. Laboratory organisms are those defined in the following list and must be contained under laboratory or animal house conditions: guinea pigs, hamsters, mice, rats, rabbits or micro-organisms. Work in all other animals and plants is not permitted.

3. For all uses in non-laboratory organisms (e.g. chickens, sheep, cattle, etc.) or plants a separate application for in vivo use must be lodged with, and approved by the Department of Agriculture. This also applies if the product is to be used in veterinary vaccine or veterinary therapeutic manufacture.

4. This Import Permit does not permit the direct or indirect exposure of the imported materials or derivatives to non-laboratory organisms or plants.

5. It is the importer's responsibility to ensure that the goods are labelled "In vitro use or in vivo use in laboratory organisms only" on the smallest packaged unit prior to distribution.

6. It is the importer's responsibility to ensure compliance with all international (e.g. IATA) and domestic requirements concerning the safe handling, transport and labelling of biological material.

7. 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.

PC6282 **Manufacturer's declaration**

Diagnostic test kits and/or research kits, not testing for microorganisms, viruses or prions

Declaration requirements

Condition	Condition Text

1. Each consignment must be accompanied by a manufacturer's declaration, stating:

a) The diagnostic kit/s and/or research kit/s does not contain any viruses, bacteria or any other microorganisms; and

b) The diagnostic kit/s and/or research kit/s does not contain any antigens derived from viruses, bacteria or any other microorganisms; and

c) The diagnostic kit/s does not test for antibodies raised against viruses, bacteria or any other microorganisms; and

d) The only animal (including human) derived materials which may be contained in the kits are:
- antibodies purified & raised against synthetic material or against antigens derived from multicellular organisms; and/or
- antigens derived from synthetic material or multicellular organisms; and/or
- laboratory reagents including animal sera, purified animal proteins, hormones, albumins (including bovine serum albumin), enzymes and lipids; and
- in volumes of no greater than 20g/20ml per individually packaged unit.

e) For reagents in individual quantities greater than 20mL or 20g, the animal (including human) derived material is no greater than 20mL or 20g per individually packaged unit.

Format of the manufacturer's declaration

1. Manufacturer
- The declaration must be made by the individual manufacturing site or the manufacturer's head office in the country of export.

2. Letterhead
- The declaration must be on manufacturer's letterhead.

3. Identification of goods.
- The declaration must be specific to the product(s) listed on this permit. Product names must be included on the manufacturers' declaration.
- The declaration must have a unique identifiable link to the consignment i.e. container number, way 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.

4. Statements
- All statements must be written in English.
- All statements must be legible and appear above the signature of the issuing officer.
- The declaration must contain the correct statement/s as required by the import conditions.
- The declaration must be free from erasures and non certified alterations. All erasures and alterations must be endorsed by the issuer of the document. Note: The only acceptable endorsement of alterations is a company seal and the signature of the company officer responsible for signing the declaration, applied adjacent to the alteration.

5. Identification of the issuing officer.
- The declaration must state the name, position and title of the issuing officer.
- Declarations must bear the signature of the issuing officer

Condition	Condition Text
	6. Date - The declaration must state the date of issue - Declarations must be issued within the last 6 months (unless otherwise specified in this import permit) General: All documentation must meet the requirements of the Department of Agriculture's minimum documentary requirements policy. For full details of the department's minimum documentary requirements, please refer to http://www.daff.gov.au/biosecurity/import/general-info/documentary-requirements.
PC6369	**Manufacturer's declaration requirements – Human Diagnostic Tests and Human Diagnostic kits** 1) Each consignment must be accompanied by a manufacturer's declaration stating that: a) The diagnostic kit(s)/test(s) are testing human conditions only; and b) All animal (including human) derived material contained in these diagnostic kit(s)/test(s) is in volumes of no greater than 20 ml or 20 g per individually packaged unit; and c) For reagents in individual quantities greater than 20mL or 20g, the animal derived material is no greater than 20mL or 20g per individually packaged unit; and d) The diagnostic kit/s do not test for pathogens and diseases listed in permit condition (PC) 6806; and e) The diagnostic kit/s do not contain any components derived from or raised against pathogens listed in permit condition (PC) 6806 [This includes, whole pathogen, parts of pathogen, antigens and antibodies raised against these pathogens]; and f) i) All components of the diagnostic kit/s derived from microorganisms, viruses, prions and fungi are incapable of replicating; or ii) The diagnostic kits/tests do not contain any components derived from microorganisms, viruses, prions or fungi. The manufacturer's declaration must be from the manufacturer of the diagnostic kits/tests. . 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. . 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

Condition	Condition Text
	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 Department of Agriculture minimum documentary requirements, please refer to http://www.daff.gov.au/biosecurity/import/general-info/documentary-requirements.
PC6760	**This import permit allows for the importation of:** 1. Transgenes (the specific gene of interest) from microorganisms and viruses, listed in PC6800, in purified cloning vectors and expression vectors i.e. bacterial plasmids, cosmid vectors, yeast artificial chromosomes, bacterial artificial chromosomes, human immunodeficiency virus (HIV) Lentivirus vectors and bacteriophages. 2. The cloning vectors may include the whole genome from any of the microorganisms and viruses listed in PC6800 3. The microorganisms and viruses listed in PC6800 may also contain cloning vectors and expression vectors i.e. bacterial plasmids, cosmid vectors, yeast artificial chromosomes, bacterial artificial chromosomes, bacteriophages and DNA inserts. These cloning vectors may be imported "empty" or may contain transgenes (the specific gene of interest) from either: i) multicellular organisms (excluding plants or fungi); or ii) from any microorganism/s and viruses listed in PC6800 • For a list of microorganisms that do not require an import permit please refer to ICON case: Microorganisms- as listed in PCT1104 • The microorganisms listed in PC6800 may also be imported on a non-biological matrix (e.g. biological indicators, spore strips).
PC6792	**Human Diagnostic Tests and Human Diagnostic kits (excluding those testing for human pathogens and diseases as listed in PC6806)**

Condition	Condition Text
	Laboratory material for in vitro use only 1. This permit allows for the importation of: a) Human diagnostic kits testing for human pathogens and diseases, excluding those listed in PC6806; and/or b) Human diagnostic kits/tests not testing for pathogens and diseases, such as haematology tests; hormone tests; drug tests; genetic tests; PCR tests etc.; and/or c) Associated reagents, standards, controls and calibrators. **Post entry / end use conditions** 2. This Import Permit allows for the importation of goods for in vitro laboratory studies only. 3. This Import Permit does not permit the use of the samples for microbiological cultures or viral isolation (except for the purpose of identifying the microorganism or virus). 4. This Import Permit does not permit the direct or indirect exposure of the imported materials or derivatives to animals (including laboratory animals) or plants. 5. For in vivo use in animals (including laboratory animals) or plants a separate application for in vivo use must be lodged with, and approved by the Department of Agriculture. 6. It is the importer's responsibility to ensure that the goods are labelled "In vitro use only" or equivalent on the smallest packaged unit prior to distribution. 7. 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. 8. 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
PC6797	**Low risk genetic material** 1. This permit allows for the importation of: a. Purified genetic material from multicellular organisms (excluding plants and fungi); and/or b. Purified cloning vectors and expression vectors i.e. bacterial plasmids, cosmid vectors, yeast artificial chromosomes, bacterial artificial chromosomes and bacteriophages may be imported "empty" or may contain transgenes (the specific gene of interest) from multicellular organisms (excluding plants or fungi) only. **Note:** · This case does not allow for cloning vectors that contain transgenes (the specific gene of

Condition	Condition Text

interest) derived from microorganisms (including viruses). For purified cloning vectors and/or live low risk microbes carrying cloning vectors containing transgenes (the specific gene of interest) derived from microorganisms (including viruses) please see ICON.

· For genetic material derived from plants please refer to ICON.

· For genetic material derived from fungi please contact Plant Programs

PC6799 **Laboratory material**

1. This permit allows for the importation of:

a) Purified & animal derived:

· Albumins, including bovine serum albumin.

· Carboxylic acids

· Co-factors

· Enzymes

· Enzyme inhibitors

· Growth factors

· Hormones

· Lipids (This includes fats, waxes, sterols, fat-soluble vitamins (e.g. A, D, E, and K), glycerides, phospholipids and their derivatives)

· Molecules (Excluding genetic material)

· Proteins (This includes derivatives e.g. peptides, amino acids). This case does not allow the import of prions.

· Vitamins

b) Fermented & then purified:

· Laboratory material derived from a fermentation process e.g. antibiotics and enzymes (It is the importers responsibility to provide documentation to support this claim)

c) Purified bacterial (including recombinant bacterial) and/or fungi derived:

Condition	Condition Text

· Antibiotics e.g. Antibiotic sensitivity discs

· Enzymes e.g. polymerases, modifying enzymes and restriction enzymes

· Growth factors

· Hormones

· Lipids (This includes fats, waxes, sterols, fat-soluble vitamins (e.g. A, D, E, and K), glycerides, phospholipids and their derivatives)

· Molecules (Excluding genetic material)

· Proteins (This includes derivatives e.g. peptides, amino acids). This case does not allow the import of prions.

Notes:

- This case does **not** allow for the import of composite products e.g. diagnostic kits. Please see "Diagnostic kits" in ICON
- The product may contain substances that are covered by the ICON case "Organic chemicals and substances – highly purified" or substances that are 100% synthetic.
- This case does **not** allow the import of antibodies. Please see "Antibodies" in ICON
- This case does **not** allow the import of live whole, whole inactivated or unpurified material derived from microorganism or prions. Please see "Microorganisms" on ICON
- An import permit is not required for purified plant derived material and alcohols, vitamins and amino acids derived from a fermentation process. The ICON case "Organic chemicals and substances – highly purified" must be met.
- An import permit is not required for yeast and its derivatives. Please see "Starter cultures" in ICON

PC6800 1. Department of Agriculture approved microorganisms (including viruses)

Achromobacter spp.	*Acidianus* spp.
Acidiphilium spp.	*Acidithiobacillus* spp.
Adeno-associated virus Types 1-9 (human AAVs only)	*Aeromonas hydrophila*
Alyciclobacillus spp.	*Amycolatopsis* spp.
Aneurinibacillus migulanus (formerly *Bacillus migulanus*)	*Aspergillus* spp.
Aquifex spp.	*Azotobacter* spp.
Bacillus atrophaeus (formerly *Bacillus subtilis* var. *niger*)	*Bacillus brevis*
Bacillus cereus	*Bacillus lichenformis*
Bacillus megaterium (excluding pv. *cerealis*)	*Bacillus pumilis* (also known as *Bacillus mesentericus* and

Condition	Condition Text

	Bacillus aminoglucosidicus
Bacillus sphaericus	*Bacillus stearothermophilus*
Bacillus subtilis	*Bacteroides* spp.
Bartonella spp.	*Bordetella* spp.
Botryococcus spp.	*Brachyspira* spp.
Brevibacillus spp.	*Burkholderia pseudomallei*
Campylobacter spp.	*Caulobacter* spp.
Chlamydophila pneumoniae (previously *Chlamydia pneumoniae)*	*Chlamydia trachomatis*
Chryseobacterium spp. (excluding *C. scophthalmum*)	*Citrobacter* spp.
Clostridium spp.	*Corynebacterium* spp. (excluding *C. pseudotuberculosis* and *C. pekinese*)
Cronobacter spp.	*Cryptococcus* spp.
Cryptomonas spp.	*Cryptosporidium* spp.
Desulfobacter spp.	*Desulfovibrio* spp.
Entamoeba spp.	*Enterobacter* spp.
Enterococcus spp.	*Enterovirus* (of human origin only and excluding *Swine Vesicular Disease Virus*)
Escherichia spp.	*Ferroplasma* spp.
Geobacillus spp.	*Geobacter* spp.
Giardia spp.	*Haemophilus* spp.
Helicobacter spp.	Human Adenovirus Types 1 – 51
Human coxsackieviruses 1 – 24	Human echovirus 1 – 33
Human herpesvirus 1 – 8 (includes Herpes simplex virus 1 & 2, *Varicella zoster*, Epstein-Barr virus and Cytomegalovirus)	Human hepatitis virus A, B, C, D, E, G & TTV
Human Immunodeficiency Virus (HIV)	Human noroviruses
Human papilloma virus	Human respiratory syncytial virus
Human rhinovirus	*Klebsiella* spp.
Legionella spp.	*Leptospira copenhageni* (*Leptospira interrogans* serovar Copenhageni)
Leptospira gripptotyphosa (*Leptospira interrogans* serovar Gripptotyphosa)	*Leptospira hardjobovis* (*Leptospira interrogans* serovar Hardjobovis)
Leptospira icterohaemorrhagiae (*Leptospira interrogans* serovar Icterohaemorrhagiae)	*Leptospira pomona* (*Leptospira interrogans* serovar Pomona)

Condition	Condition Text

Leptospirillium spp.	*Listeria* spp.
Magnetospirillium spp. (formerly *Aquaspirillium* spp.)	Metapneumonovirus (Human)
Metarhizium anisopliae	*Methanococcus* spp.
Moraxella spp. (includes subgen. *Branhamella* and subgen. *Moraxella*) (excluding *M. anatipestifer*)	*Morganella* spp.
Murine Cytomegalovirus (MCMV)	Murine leukaemia virus
Mycobacterium spp. (excluding *M. bovis* and *M. caprae*)	*Mycoplasma pneumoniae*
Neisseria spp.	*Nippostrongylus brasiliensis*
Parainfluenza virus (human)	*Pediococcus* spp.
Penicillium chrysogenum	*Porphyromonas* spp.
Proteus spp.	*Providencia* spp.
Pseudomonas aeruginosa	*Pseudomonas fluorescens* (excluding biovar II)
Pseudomonas putida	*Rhodobacter* spp.
Rhodococcus spp.	*Roseomonas* spp.
Rubrivivax spp.	*Saccharopolyspora* spp.
Salmonella adelaide	*Salmonella agona* (*Salmonella enterica* subsp. *enterica* serovar Agona)
Salmonella derby (*Salmonella enterica* subsp. *enterica* serovar Derby)	*Salmonella salford*
Salmonella seftenburg (*seftenberg*)	*Serratia* spp.
Shewanella spp.	*Shigella* spp.
Sindbis virus	*Staphylococcus* spp.
Stentrophomonas spp.	*Streptococcus* spp.
Sulfobacillus spp.	*Sulfolobus* spp.
Sulfurisphaera spp.	*Tetrahymena* spp.
Thermus spp.	*Thiobacillus* spp.
Toxoplasma spp.	Vaccinia virus (cow pox)
Vibrio alginolyticus	*Vibrio cholerae* (excluding serotype 01 and serotype 0139)
Vibrio parahaemolyticus	*Vibrio vulnificus* (excluding biovar II)

PC6805 **Genetic material, DNA, cDNA and RNA derived from the following micro-organisms and viruses may NOT be imported using this import permit:**

1) All plant pathogens (viruses, viroids, bacteria, fungi and stramenopiles)

2) Micro-organisms associated with Quarantinable diseases of humans listed in Table 9 of

Condition	Condition Text

Quarantine Proclamation 1998

· Rabies (Lyssavirus)

· Severe Acute Respiratory Syndrome (SARS) (SARS associated Coronavirus)

· Smallpox (Variola virus and Poxvirus variola)

· Viral haemorrhagic fevers of humans including Ebola haemorrhagic fever (Filoviridae), Marburg virus (Filoviridae), Lassa Fever (Arenaviridae) and Crimean-Congo hemorrhagic fever (Nairovirus)

· Yellow fever (Flavivirus)

· Highly Pathogenic Avian Influenza in Humans

3) Foot and mouth disease virus

4) Rinderpest virus

5) African horse sickness virus

6) Peste des petits ruminants virus

7) Ovine and caprine pox virus

8) Pulmonary adenomatosis virus

9) Swine vesicular disease virus

10) African swine fever virus

11) Classical swine fever virus

12) Avian influenza virus

13) Newcastle disease virus

PC6806 **Diagnostic kits, reagents and standards containing or testing for the following human pathogens and diseases (or diseases with zoonotic potential) may NOT be imported using this import permit.**

Micro-organisms associated with Quarantinable diseases of humans listed in Table 9 of *Quarantine Proclamation 1998*

· Cholera (Vibriocholerae)

Condition	Condition Text

· Highly Pathogenic Avian Influenza in Humans

· Plague (Yersiniapestis)

· Rabies (Lyssavirus)

· Severe Acute Respiratory Syndrome (SARS) (severe acute respiratory syndrome-related coronavirus)

· Smallpox (Variola virus and Poxvirus variola)

· Viral haemorrhagic fevers of humans including Ebola haemorrhagic fever (Filoviridae), Marburg virus (Filoviridae), Lassa Fever (Arenaviridae) and Crimean-Congo hemorrhagic fever (Nairovirus)

· Yellow fever (Flavivirus)

Diagnostic kits, reagents and standards containing or testing for the following pathogens and diseases with zoonotic potential may NOT be imported using this import permit

Viruses and Prions:

· Avian influenza (avian influenza virus)

· Equine encephalomyelitis (Eastern, Venezuelan and Western equine encephalomyelitis viruses)

· Foot-and-mouth disease (foot-and-mouth disease virus)

· Japanese encephalitis (Japanese encephalitis virus)

. Hantaan virus (Korean haemorrhagic fever virus)

· Louping ill (also known as Russian spring summer encephalitis, Central European encephalitis, caused by Louping ill virus)

. Lymphocytic choriomeningitis (Lymphocytic choriomeningitis virus)

· Newcastle disease (Newcastle disease virus)

· Nipah virus encephalitis (Nipah virus)

· Rift Valley fever (Rift Valley fever virus)

· St Louis Encephalitis (St Louis Encephalitis virus)

· Swine influenza (swine influenza virus)

· Transmissible spongiform encephalopathy (Bovine spongiform encephalopathy, scrapie, variant Creutzfeldt-Jakob disease, prion protein)

Condition	Condition Text

· Vesicular stomatitis (Vesicular stomatitis virus)

· West Nile fever (West Nile virus)

Bacteria and fungi:

· Anthrax *(Bacillus anthracis)*

· Bovine tuberculosis (*Mycobacterium bovis)*

· Brucellosis (*Brucellaabortus, B. canis, B.melitensis*)

· Enzootic abortion of ewes (*Chlamydophilaabortus*)

· Epizootic lymphangitis or histoplamosis (*Histoplasmacapsulatum* var. *farciminosum)*

· Glanders (*Burkholderiamallei*, formerly *Pseudomonas mallei*)

· Salmonellosis (*Salmonella* Abortusovis, *Salmonella* Enteritidis, *Salmonella* Gallinarum, *Salmonella Pullorum*)

· Tularemia (*Francisellatularensis)*

PC6855 **This condition allows for the importation of genetic material purified & derived from microorganisms & viruses (excluding those listed in PC6805).**

Labelling requirement

The goods must be clearly labelled with the name of the source microorganism or virus.

PC6856 **Manufacturer's declaration requirements – Genetic material - purified & derived from microorganisms & viruses (excluding high risk species)**

1) Each consignment must be accompanied by a manufacturer's declaration stating that:

The genetic material has been highly purified and is unable to replicate.

The manufacturer's declaration must be from the manufacturer of the genetic material.

. 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.
. written in English.
. signed by a designated representative whose name, position and title also appear.
. identify the date of issue.

Condition	Condition Text

. 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 Department of Agriculture minimum documentary requirements, please refer to http://www.daff.gov.au/biosecurity/import/general-info/documentary-requirements.

End of Condition Text

Delegate of Director of Quarantine
Printed Name Tran Tang **Date** 28 Jul 2015