Australian Government
Department of Agriculture, Water and the Environment

Permit to import conditionally non-prohibited goods

This permit is issued under *Biosecurity Act 2015* Section 179 (1)

Permit: 0006011256

Valid for: multiple consignments between 20 February 2022 and 20 February 2024

This permit is issued to: Diversity Arrays Technology Pty Ltd
Building 3 Level D
University of Canberra
Kirinari Street
BRUCE ACT 2617
Australia

Attention: Ms Bianca Mauch

This permit is issued for the import of Biological products (Standard goods).

Exporter details:	Various exporters

This permit includes the following good(s). Refer to the indicated page for details of the permit conditions:

1. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excluding reproductive material) from species, other than those excluded	Page 5
2. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excl. viable reproductive material) sourced from captive primates only	Page 7
3. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	

This permit is granted subject to the requirement that fees determined under section 592(1) are paid.

Matthew Byatt
Delegate of the Director of Biosecurity

Date: 21 January 2022

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Permit Conditions:	Animal fluids and tissues (excluding reproductive material) sourced from bovines only	Page 10

4. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excluding reproductive material) sourced from ovines and caprines only	Page 13

5. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excluding reproductive material) sourced from cervines (deer) only	Page 16

6. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excluding reproductive material) sourced from camelids only	Page 19

7. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excluding reproductive material) sourced from suids (porcines) only	Page 22

8. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excl. reproductive material) sourced from equines only	Page 25

9. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excl. reproductive material) sourced from equines, containment required	Page 28

10. Animal fluids and tissues (excl. viable reproductive material)		
End use:	In vitro use or in vivo use in laboratory organisms	
Country of export:	Various countries	
Country of origin:	Various countries	
Permit Conditions:	Animal fluids and tissues (excluding reproductive material) sourced from avians only	Page 31

NOTE: Where a good has more than one set of permit conditions please read each set to determine which set of permit conditions applies to a specific consignment.

---------- End of commodity list ----------

Important information about this permit and the import of goods

Note: This permit covers Department of Agriculture, Water and the Environment import conditions. It is the permit holder's responsibility to ensure all legal requirements relating to the goods described in this permit are met. While the permit holder should rely on their own inquiries, the following information is provided to assist the permit holder in meeting legal obligations in relation to the importation of the goods described in this permit.

Information about this permit

Authority to import

The permit holder is authorised to import the goods described in this permit subject to the listed conditions specified in this permit.

Compliance with permit conditions and assessment and management of biosecurity risk

All imports are subject to biosecurity control and may be subject to biosecurity inspection on arrival to determine compliance with the listed permit conditions and to assess the level of biosecurity risk associated with the goods. Imports that do not comply with the import conditions specified in the permit may present an unacceptable level of biosecurity risk and may be subject to biosecurity measures that may include treatment, export or destruction at the permit holder's expense or forfeited to the Commonwealth.

Additionally, non-compliance with import permit conditions may constitute an offence or contravention of a civil penalty provision under section 187 of the *Biosecurity Act 2015*.

Change of import conditions

The Director of Biosecurity may, in accordance with section 180 of the *Biosecurity Act 2015* vary or revoke the conditions on a permit or impose further conditions.

General information about importing goods

Notification of import

Notification of the import must be provided to the Department of Agriculture, Water and the Environment 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,* or where other exceptions specified in the *Biosecurity Regulation 2016* apply. Notification must be provided in accordance with section 120 of the *Biosecurity Act 2015* and Part 1 of Chapter 2 of the *Biosecurity Regulation 2016*. Please refer to 'Sending your goods to Australia' on the Department of Agriculture, Water and the Environment website.

Provision of required documentation

It is recommended that all required documentation accompanies each consignment. Required documentation must be presented to the Department of Agriculture, Water and the Environment for assessment. Airfreight or mail shipments should have all required documentation securely attached to the outside of the package, and clearly marked "Attention Department of Agriculture, Water and the Environment". Documentation may include the permit (or permit number), government certification and invoice.

If the product description on the permit varies from the identifying documentation provided, the goods will not be released from biosecurity control unless evidence is provided to the biosecurity officer that the permit covers the goods in the consignment.

Any documentation provided must comply with the Department of Agriculture, Water and the Environment's minimum documentation requirements policy.

Non-commodity cargo clearance

In addition to the conditions for the goods being imported, non-commodity biosecurity risks are assessed including container cleanliness, packaging and destination concerns, and may be subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

Fees

Fees are payable to the Department of Agriculture, Water and the Environment for certain services (see the *Biosecurity Charges Imposition (General) Regulation 2016*, Part 2 of Chapter 9 of the *Biosecurity Regulation 2016* and Part 3 of Chapter 11 of the *Biosecurity Act 2015*). Detail on how the department applies fees and levies may be found in the Charging guidelines.

Compliance with other regulatory provisions

Goods imported into Australia may be subject to regulatory requirements under other legislation. It is the permit holder's responsibility to identify and ensure they have complied with all requirements of any other regulatory agency or advisory body prior to and after importation.

Permit conditions

It is the importer's responsibility to ensure that the following permit conditions are met in relation to each consignment. Where more than one set of permit conditions is shown for a good please read each set of conditions to determine which applies to a specific consignment.

1. Animal fluids and tissues (excluding reproductive material) from species, other than those excluded

This section contains permit conditions for the following commodity (or commodities):

1.	Animal fluids and tissues (excl. viable reproductive material)

1.1. Biosecurity Pathway

a. **Sourcing**
The goods must be animal fluids and tissues only.

The goods must not be reproductive material.

b. The goods must not be sourced from: avians, bovines, camelids, caprines, cervines, equines, giraffids, ovines, prawns, primates, suids (porcines) or Salmonidae fish.

c. **Animal Health**
The goods must not be sourced from animals with signs of infectious disease at the time of collection.

The goods must not have been deliberately infected with a disease agent other than those listed below.

Antisera may only be raised against:

1. synthetic material, or
2. antigens derived from multicellular organisms, or
3. starter cultures (Appendix 1), or
4. standard laboratory microorganisms (including viruses) list (Appendix 2).

d. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

e. **Packaging**
The goods must be imported in quantities of no greater than:

1. 20mL or 20g for each individually packaged unit, or
2. for urine only, 500mL or 500g for each individually packaged unit.

f. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

g. **Commercial administrative conditions**
Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).
   e.g. 1: Product XRab = Purified protein derived from rabbits
   e.g. 2: Product AX = Synthetic antibiotic
   e.g. 3: Comte = Cheese.

h. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

i. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

2. Animal fluids and tissues (excl. viable reproductive material) sourced from captive primates only

This section contains permit conditions for the following commodity (or commodities):

2.	Animal fluids and tissues (excl. viable reproductive material)

2.1. Biosecurity Pathway

a. **Source species**

The goods must be fluids and tissues sourced from captive primates that are held within laboratory or zoological facilities only.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods were obtained from primates held within a laboratory or zoological facility only.
2. A statement that the goods:

2.1. are not reproductive material, or

2.2. the reproductive material is:
2.2.1. non-viable,

2.2.2. is transported at room temperature, and

2.2.3. is not intended for use in artificial insemination (AI) or assisted reproductive treatment (ART).

[The declaration must indicate the option that applies.]

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

3.1. the goods are not antisera, or

3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

[The declaration must indicate the option that applies].

c. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

d. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

e. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

f. **Commercial administrative conditions**

Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).

   e.g. 1: Product XRab = Purified protein derived from rabbits

   e.g. 2: Product AX = Synthetic antibiotic

   e.g. 3: Comte = Cheese.

g. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture,

Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

h. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

3. Animal fluids and tissues (excluding reproductive material) sourced from bovines only

This section contains permit conditions for the following commodity (or commodities):

3.	Animal fluids and tissues (excl. viable reproductive material)

3.1. Biosecurity Pathway

a. **Source species and countries**

The goods must be fluids and tissues sourced from bovines only, which resided in countries approved for bovine fluids and tissues (as listed on the department's website) at the time of collection.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods are of <<*insert species of animal*>> origin only.
2. A statement that the goods have only been sourced from animal/s residing in <<*insert name/s of country/ies*>>.
3. A statement that the goods are not reproductive material.

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

   3.1. the goods are not antisera, or

   3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

   [The declaration must indicate the option that applies].

c. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

d. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

e. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

f. **Commercial administrative conditions**

Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).

   e.g. 1: Product XRab = Purified protein derived from rabbits

   e.g. 2: Product AX = Synthetic antibiotic

   e.g. 3: Comte = Cheese.

g. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

h. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be

subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

4. Animal fluids and tissues (excluding reproductive material) sourced from ovines and caprines only

This section contains permit conditions for the following commodity (or commodities):

4.	Animal fluids and tissues (excl. viable reproductive material)

4.1. Biosecurity Pathway

a. **Source species and countries**

The goods must be fluids and tissues sourced from ovines and/or caprines only, which resided in countries approved for ovine and caprine fluids and tissues (as listed on the department's website) at the time of collection.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods are of <<*insert species of animal*>> origin only.
2. A statement that the goods have only been sourced from animal/s residing in <<*insert name/s of country/ies*>>.
3. A statement that the goods are not reproductive material.

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

   3.1. the goods are not antisera, or

   3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

   [The declaration must indicate the option that applies].

c. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

d. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

e. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

f. **Commercial administrative conditions**

Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).

   e.g. 1: Product XRab = Purified protein derived from rabbits

   e.g. 2: Product AX = Synthetic antibiotic

   e.g. 3: Comte = Cheese.

g. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

h. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be

subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

5. Animal fluids and tissues (excluding reproductive material) sourced from cervines (deer) only

This section contains permit conditions for the following commodity (or commodities):

5.	Animal fluids and tissues (excl. viable reproductive material)

5.1. Biosecurity Pathway

a. **Source species and countries**

The goods must be fluids and tissues sourced from cervines only, which resided in countries approved for cervine fluids and tissues (as listed on the department's website) at the time of collection.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods are of <<*insert species of animal*>> origin only.
2. A statement that the goods have only been sourced from animal/s residing in <<*insert name/s of country/ies*>>.
3. A statement that the goods are not reproductive material.

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

   3.1. the goods are not antisera, or

   3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

   [The declaration must indicate the option that applies].

c. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

d. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

e. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

f. **Commercial administrative conditions**

Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).

   e.g. 1: Product XRab = Purified protein derived from rabbits

   e.g. 2: Product AX = Synthetic antibiotic

   e.g. 3: Comte = Cheese.

g. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

h. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be

subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

6. Animal fluids and tissues (excluding reproductive material) sourced from camelids only

This section contains permit conditions for the following commodity (or commodities):

6.	Animal fluids and tissues (excl. viable reproductive material)

6.1. Biosecurity Pathway

a. **Source species and countries**

The goods must be fluids and tissues sourced from camelids only, which resided in countries approved for camelid fluids and tissues (as listed on the department's website) at the time of collection.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods are of <<*insert species of animal*>> origin only.
2. A statement that the goods have only been sourced from animal/s residing in <<*insert name/s of country/ies*>>.
3. A statement that the goods are not reproductive material.

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

   3.1. the goods are not antisera, or

   3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

   [The declaration must indicate the option that applies].

c. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

d. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

e. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

f. **Commercial administrative conditions**

Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).

   e.g. 1: Product XRab = Purified protein derived from rabbits

   e.g. 2: Product AX = Synthetic antibiotic

   e.g. 3: Comte = Cheese.

g. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

h. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be

subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

7. Animal fluids and tissues (excluding reproductive material) sourced from suids (porcines) only

This section contains permit conditions for the following commodity (or commodities):

7.	Animal fluids and tissues (excl. viable reproductive material)

7.1. Biosecurity Pathway

a. **Source species and countries**

The goods must be fluids and tissues sourced from suids (porcines) only, which resided in countries approved for suid fluids and tissues (as listed on the department's website) at the time of collection.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods are of <<*insert species of animal*>> origin only.
2. A statement that the goods have only been sourced from animal/s residing in <<*insert name/s of country/ies*>>.
3. A statement that the goods are not reproductive material.

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

   3.1. the goods are not antisera, or

   3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

   [The declaration must indicate the option that applies].

c. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

d. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

e. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

f. **Commercial administrative conditions**

Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).

   e.g. 1: Product XRab = Purified protein derived from rabbits

   e.g. 2: Product AX = Synthetic antibiotic

   e.g. 3: Comte = Cheese.

g. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

h. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be

subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

8. Animal fluids and tissues (excl. reproductive material) sourced from equines only

This section contains permit conditions for the following commodity (or commodities):

8.	Animal fluids and tissues (excl. viable reproductive material)

8.1. Biosecurity Pathway

a. **Source species and countries**

The goods must be fluids and tissues sourced from equines only, which resided in countries approved for equine fluids and tissues (as listed on the department's website) at the time of collection.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods are of <<*insert species of animal*>> origin only.
2. A statement that the goods have only been sourced from animal/s residing in <<*insert name/s of country/ies*>>.
3. A statement that the goods are not reproductive material.

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

   3.1. the goods are not antisera, or

   3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

   [The declaration must indicate the option that applies].

c. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

d. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

e. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

f. **Commercial administrative conditions**

Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).

   e.g. 1: Product XRab = Purified protein derived from rabbits

   e.g. 2: Product AX = Synthetic antibiotic

   e.g. 3: Comte = Cheese.

g. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

h. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be

subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

9. Animal fluids and tissues (excl. reproductive material) sourced from equines, containment required

This section contains permit conditions for the following commodity (or commodities):

9.	Animal fluids and tissues (excl. viable reproductive material)

9.1. Biosecurity Pathway

a. **Source species and countries**

The goods must be fluids and tissues sourced from equines only, which resided in countries approved for equine fluids and tissues with containment (as listed on the department's website) at the time of collection.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods are of <<*insert species of animal*>> origin only.
2. A statement that the goods have only been sourced from animal/s residing in <<*insert name/s of country/ies*>>.
3. A statement that the goods are not reproductive material.

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

   3.1. the goods are not antisera, or

   3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

   [The declaration must indicate the option that applies].

c. The goods are for use at an approved arrangement site class 5. The level of containment must be BC level 1 or higher.

These approved arrangement site/s must have current approval from the Department of Agriculture, Water and the Environment as a class 5 approved arrangement site/s at the time of importation and until such time that all imported material and its derivatives are removed for disposal or export.

d. If the above conditions cannot be met, or the goods cannot be directed to an appropriate approved arrangement site, the goods must be treated with ionising irradiation to a level that achieves a minimum absorbed dose of 50kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

e. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

f. **Post entry/end use conditions**

Approved end use:

1. *in vitro* laboratory studies.

The following end uses are not permitted:

1. in culturing or isolating microorganisms and infectious agents,
2. in the synthesis of replication-competent microorganisms, infectious agents or homologues.

It is the importer's responsibility to ensure that the goods are labelled "*in vitro* only" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions,
2. *in vivo* in non-laboratory organisms e.g. chickens, sheep, cattle,
3. as veterinary vaccines and therapeutics,
4. in plants.

For more information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090.

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

g. In addition to the standards for waste disposal outlined in the 5.1 AA standards and ASNZ 2243.3, liquid biosecurity waste which is or has come into contact with imported material in this category must be decontaminated prior to disposal as biosecurity waste by one of the following disinfectant methods:

1. Virkon – final concentration of 10 g per 1 L for at least 10 minutes or as per the manufacturer's instructions.
2. Chlorine (i.e. sodium hypochlorite solutions) – final concentration of 1% (10,000ppm

available chlorine) for a minimum of 10 minutes.

h. **Commercial administrative conditions**
Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).
   e.g. 1: Product XRab = Purified protein derived from rabbits
   e.g. 2: Product AX = Synthetic antibiotic
   e.g. 3: Comte = Cheese.

i. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

j. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

10. Animal fluids and tissues (excluding reproductive material) sourced from avians only

This section contains permit conditions for the following commodity (or commodities):

10. Animal fluids and tissues (excl. viable reproductive material)

10.1. Biosecurity Pathway

a. **Source species and countries**

The goods must be fluids and tissues sourced from avians only, which resided in countries approved for avian fluids and tissues (as listed on the department's website) at the time of collection.

b. The goods must meet biosecurity requirements.

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

i. **Sourcing**

1. A statement that the goods are of <<*insert species of animal*>> origin only.
2. A statement that the goods have only been sourced from animal/s residing in <<*insert name/s of country/ies*>>.
3. A statement that the goods are not reproductive material.

AND

ii. **Animal Health**

1. A statement that the goods were sourced from animals with no clinical signs of infectious disease at the time of collection.
2. A statement that the goods have not been deliberately infected with a disease agent.
3. A statement that either:

   3.1. the goods are not antisera, or

   3.2. the antisera has been raised against synthetic material or against antigens derived from multicellular organisms.

   [The declaration must indicate the option that applies].

c. If the above conditions cannot be met, the goods must be treated with ionising radiation to a level that achieves a minimum absorbed dose of 50 kGy before being released to the importer. Irradiation on arrival is mandatory, even if the goods have been treated prior to import.

d. The goods must meet biosecurity requirements

To demonstrate compliance with this requirement you must present the following on a Manufacturer's declaration or Supplier's declaration:

Packaging

A statement that the goods are either:

1. individually packaged in units of no greater than 20mL or 20g, or
2. urine only, and are individually packaged in units no greater than 500mL.

[The declaration must indicate the option that applies].

e. **Post entry/end use conditions**

Approved end uses:

1. *in vitro* laboratory studies, and/or
2. *in vivo* in laboratory organisms. Laboratory organisms are guinea pigs, hamsters, mice, rats, rabbits or microorganisms contained under laboratory or animal house conditions.

These conditions do not permit:

1. culturing or isolating microorganisms and infectious agent.
2. the synthesis of replication-competent microorganisms, infectious agent or homologues.

It is the importers responsibility to ensure that the goods are labelled "*in-vitro or in-vivo use in laboratory organisms only*" on the smallest packaged unit, prior to distribution. The products may be labelled post entry.

Additional written approvals are required prior to direct or indirect use:

1. in non-laboratory organisms e.g. chickens, sheep, cattle.
2. in plants.

For information on how to obtain additional written approvals contact imports@awe.gov.au or call 1800 900 090

Where applicable, the importer or end user must comply with:

1. International (e.g. International Air Transport Association) and domestic requirements concerning the safe handling, transport and labelling of biological material
2. AS/NZS 2243 Safety in Laboratories standards
3. Office of the Gene Technology Regulator (OGTR) requirements
4. The Security Sensitive Biological Agents (SSBA) regulatory scheme.

f. **Commercial administrative conditions**

Documents must be provided with each consignment which:

1. identify the consignment (if non-personal) e.g. entry number
2. identify all goods being imported as part of this consignment e.g. invoice or waybill or importer's manifest
3. describe the goods being imported (where not clear).

   e.g. 1: Product XRab = Purified protein derived from rabbits

   e.g. 2: Product AX = Synthetic antibiotic

   e.g. 3: Comte = Cheese.

g. Under the Biosecurity Charges Imposition (General) Regulation 2016 and Chapter 9, Part 2 of the Biosecurity Regulation 2016, fees are payable to the Department of Agriculture, Water and the Environment for all services. Detail on how the department applies fees and levies may be found in the Charging guidelines.

h. In addition to the conditions for the goods being imported, non-commodity concerns must be assessed including container cleanliness, packaging and destination concerns, and may be

subject to inspection and treatment on arrival. Please refer to the Non-Commodity Cargo Clearance BICON case for further information.

Appendix 1: List: Approved starter cultures

List of approved starter cultures

Acetobacter spp.	*Aspergillus brasiliensis*	*Aspergillus oryzae*
Aspergillus niger	*Bacillus acidopullulyticus*	*Bacillus amyloliquefaciens*
Bacillus coagulans	*Bacillus halodurans*	*Bacillus licheniformis*
Bacillus subtilis	Baker's yeast	*Bifidobacterium* spp.
Brevibacterium linens	Brewer's yeast	*Candida* spp.
Chaetomium gracile	*Citeromyces* spp.	*Clavispora* spp.
Debaryomyces spp.	*Dekkera* spp.	*Enterococcus durans*
Enterococcus faecalis	*Enterococcus faecium*	*Geotrichum candidum*
Hansenula spp.	*Hasegawaea* spp.	*Humicola insolens*
Hyphopichia spp.	*Issatchenkia* spp.	*Kluyveromyces* spp.
Lactic acid bacteria	*Lactobacillus* spp.	*Lactococcus* spp.
Leuconostoc spp. (*Oenococcus* spp.)	*Monascus* spp.	*Pediococcus pentosaceus*
Penicillium camemberti (also known as *Penicillium camembertii* and *Penicillium candidum*)	*Penicillium funiculosum*	*Penicillium roqueforti* (also known as *Penicillium roquefortii*)
Phaffia spp.	*Pichia* spp.	*Propionibacterium* spp.
Rhizopus spp.	*Saccharomyces* spp.	*Schizosaccharomyces* spp.
Schwanniomyces spp.	*Staphylococcus carnosus*	*Staphylococcus xylosus*
Streptococcus cremoris	*Streptococcus diacetilactis*	*Streptococcus durans*
Streptococcus faecalis	*Streptococcus lactis*	*Streptococcus salivarius*
Streptococcus thermophilus	*Streptomyces olivaceus*	*Streptomyces olivochromogenes*
Streptomyces murinus	*Streptomyces mobaraensis* (former name *Streptoverticillium mobaraensis*)	*Streptomyces rubiginosus*
Streptomyces violaceoruber	*Talaromyces emersonii (*former name *Penicillium emersonii*)	*Torulaspora* spp.
Torulopsis spp.	*Trichoderma harzianum*	*Trichoderma reesei* (former name *Trichoderma longibrachiatum*)
Trichoderma viride	Wine culture	Yoghurt/Kefir culture
Zygoascus spp.	*Zygosaccharomyces* spp.	

Appendix 2: List: Standard laboratory microorganisms and infectious agents

The following list contains microorganism and infectious agent that do not require biosecurity containment. These microorganisms are endemic (occur in Australia) and are commonly imported by laboratories in Australia.

Achromobacter spp.	*Acidianus* spp.	*Acidiphilium* spp.	*Acidithiobacillus* spp.
Acremonium cellulolyticus	*Actinomadura malachitica*	*Actinomadura viridis*	*Actinomyces rectiverticillatus*
Adeno-associated virus	*Aeromonas hydrophila*	*Alcaligenes denitrificans*	*Alicyclobacillus* spp.
Ampelomyces quisqualis	*Anabaena cylindrica*	*Anaerobacter polyendosporus*	*Aneurinibacillus migulanus (formerly Bacillus migulanus)*
Aquifex spp.	*Arthrobacter picolinophilus*	*Arthrobacter* spp.	*Aspergillus* spp.
Azorhizobium caulinodans	*Azotobacter* spp.	*Bacillus aminoglucosidicus*	*Bacillus atrophaeus (formerly Bacillus subtilis var. niger)*
Bacillus brevis syn. Brevibacillus brevis	*Bacillus cereus excluding Biovar anthracis*	*Bacillus fluorescens putidus*	*Bacillus geniculatus*
Bacillus ginsengihumi	*Bacillus licheniformis*	*Bacillus megaterium (excluding pv. cerealis)*	*Bacillus mesentericus*
Bacillus methylotrophicus	*Bacillus mojavensis*	*Bacillus pasteurii*	*Bacillus pumilus syn. Bacillus mesentericus, Bacillus aminoglucosidicus*
Bacillus putidus	*Bacillus simplex*	*Bacillus sphaericus*	*Bacillus stearothermophilus*
Bacillus subtilis	*Bacillus thuringiensis*	*Bacteroides* spp.	*Bartonella* spp.
Beauveria bassiana	*Bordetella* spp.	*Botryococcus* spp.	*Brachyspira* spp.
Brevibacillus spp. *(excluding B. laterosporus)*	*Burkholderia pseudomallei*	*Campylobacter* spp.	*Caulobacter* spp.
Chlamydia trachomatis	*Chlamydophila pneumonia*	*Chlorella* spp.	*Chryseobacterium* spp. *(excluding C. scophthalmum)*
Cicinnobolus cesatti	*Citrobacter* spp.	*Clostridium* spp.	*Comamonas acidovorans*
Corynebacterium spp. *(excluding C. pseudotuberculosis)*	*Cronobacter* spp.	*Cryptococcus* spp.	*Cryptomonas* spp.

Cryptosporidium spp.	*Dehalobacter* spp.	*Dehalococcoides* spp.	*Dehalogenimonas* spp.
Delftia acidovorans	*Desulfobacter* spp.	*Desulfovibrio* spp.	*Ensifer adhaerens*
Ensifer meliloti	*Entamoeba* spp.	*Enterobacter asburiae*	*Enterobacter* spp.
Enterococcus spp.	*Enterovirus (human origin only, and excluding swine vesicular disease virus and human enterovirus C)*	*Entomophthora anisopliae*	*Erwinia tasmaniensis*
Escherichia spp.	*Ferroplasma* spp.	*Fusarium venenatum*	*Geobacillus* spp.
Geobacter spp.	*Giardia* spp.	*Gigaspora margarita*	*Gliocadium catenalatum*
Haemophilus spp.	*Human Adenovirus Types 1-51*	*Human coxsackieviruses 1-24*	*Human echovirus 1-33*
Human hepatitis virus A, B, C, D, E, G &TTV	*Human Herpes virus 1-8 (includes Herpes simplex virus 1 and 2, Varicella zoster, Epstein-Barr virus and Cytomegalovirus)*	*Human immunodeficiency virus (HIV)*	*Human noroviruses*
Human papilloma virus	*Human respiratory syncytial virus*	*Human rhinovirus*	*Isochrysis galbana*
Klebsiella spp.	*Legionella* spp.	*Leptospira copenhageni (Leptospira interrogans serovar Copenhageni)*	*Leptospira gripptotyphosa (Leptospira interrogans serovar Gripptotyphosa)*
Leptospira hardjobovis (Leptospira borgpetersenii serovar hardjo-bovis)	*Leptospira icterohaemorrhagiae (Leptospira interrogans serovar Icterohaemorrhagiae)*	*Leptospira pomona (Leptospira interrogans serovar Pomona)*	*Leptospirillum* spp.
Listeria spp.	*Magnetospirillum* spp. *(formerly Aquaspirillum* spp.*)*	*Metapneumovirus (human)*	*Metarhizium acridum*
Metarhizium anisopliae var. anisopliae	*Methanococcus* spp.	*Microtetraspora viridis*	*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	*Nannochloropsis* spp.	*Neisseria* spp.	*Nippostrongylus*

			brasiliensis
Nocardia calcarea	*Ochrobactrum anthropi*	*Paenarthrobacter* spp.	*Paenibacillus alvei*
Paenibacillus brasiliensis	*Parainfluenza virus (human)*	*Pediococcus* spp.	*Penicillium chrysogenum*
Penicillium oxalicum	*Penicillium velutinum*	*Pleomorphomonas oryzae*	*Porphyromonas* spp.
Pristionchus americanus	*Pristionchus maupasi*	*Pristionchus pacificus*	*Proteus* spp.
Providencia spp.	*Pseudomonas acidovorans*	*Pseudomonas aeruginosa*	*Pseudomonas antarctica*
Pseudomonas citronellolis	*Pseudomonas convexa*	*Pseudomonas eisenbergii*	*Pseudomonas fluorescens (excluding biovar II)*
Pseudomonas geniculata	*Pseudomonas incognita*	*Pseudomonas monteilii*	*Pseudomonas ovalis*
Pseudomonas putida	*Pseudomonas rugosa*	*Pseudomonas striata*	*Rhabditis myriophila*
Rhizobium meliloti	*Rhodobacter* spp.	*Rhodococcus* spp.	*Roseomonas* spp.
Rubella virus	*Rubrivivax* spp.	*Saccharopolyspora spinosa*	*Saccharopolyspora* spp.
Salmonella Adelaide (Salmonella enterica subsp. enterica serovar Adelaide)	*Salmonella Agona (Salmonella enterica subsp. enterica serovar Agona)*	*Salmonella Derby (Salmonella enterica subsp. enterica serovar Derby)*	*Salmonella Salford (Salmonella enterica subsp. enterica serovar Salford)*
Salmonella Senftenburg (Salmonella enterica subsp. enterica serovar Senftenberg)	*Scutellospora dipurpurescens*	*Serratia* spp.	*Shewanella* spp. *(excluding Shewanella marisflavi)*
Shigella spp.	*Sindbis virus*	*Sinorhizobium adhaerens*	*Sinorhizobium meliloti*
Sporosarcina pasteurii	*Staphylococcus* spp.	*Stenotrophomonas* spp.	*Streptococcus* spp.
Streptomyces rectiverticillatus	*Streptoverticillium rectiverticillatum*	*Suillus granulatus*	*Sulfobacillus* spp.
Sulfolobus spp.	*Sulfurisphaera* spp.	*Tetrahymena* spp.	*Thermus* spp.
Thiobacillus spp.	*Toxoplasma* spp.	*Tritirachium shiotae*	*Tritirachium shiotae*
Vaccinia virus (cow pox)	*Vibrio alginolyticus*	*Vibrio cholerae (excluding serotype 01 and serotype 0139)*	*Vibrio parahaemolyticus (excluding VPAHPND strains with plasmid coding for Pir toxin homologues)*
Vibrio vulnificus (excluding biovar II)	*Wolinella succinogens*	*Xanthobacter* spp.	*Yersinia enterocolitica*

---------- End of permit conditions ----------