

Permit to Import Quarantine Material

Permit: **IP07002747** Valid From: **02 Feb 2007** Valid To: **02 Feb 2009** Page 1 of 3

| Importer | Exporter |
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| Australian Bioanalytical Services Pty ltd Loading Bay 5, Building 1 Princess Alexandra Hospital Ipswich Road Woolloongabba QLD 4102 Attn: Paul Salm | Various Suppliers Exporters Various Addresses In All countries |

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

Note: This permit covers AQIS quarantine requirement only.

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

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

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

It is the importer's responsibility to identify, and to ensure it has complied with, all requirements of any other regulatory organisations and advisory bodies prior to and after importation including The Australian Customs Service, The Department of Health and Ageing, Therapeutic Goods Administration, Australian Pesticides and Veterinary Medicines Authority, Department of Environment and Heritage, 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.

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

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


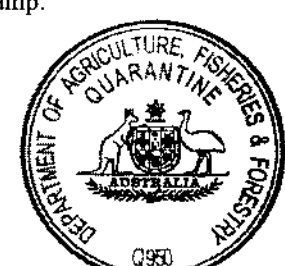
| Commodity Name | Condition Number(s) | Country | End Use |
|---|---------------------|---------------|----------|
| Human Products (Human tissues and fluids not known to be infectious) | PC4206 | All countries | In-vitro |

| Condition | Condition Text |
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PC4206

1. A valid copy of this Import Permit (or a method of identifying the Import Permit such as the Import Permit number) and all required documentation must accompany each consignment. Alternatively, necessary documentation will need to be presented to AQIS at the time of clearance. In

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

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|  Delegate of Director of Quarantine Printed Name Sally Grimes | Stamp:  |
| Date 12 Feb 2007 | |

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order to facilitate clearance, airfreight or mail shipments should have all documentation securely attached to the outside of the package, and clearly marked "Attention Quarantine". Documentation may include Import Permit (or Import Permit number), and invoice.

DOCUMENTATION REQUIREMENTS

2. Each consignment must be clearly identified and linked to the relevant item(s) on the Import Permit. Identifying documentation must be available to the quarantine officer at the time of clearance. This documentation may include:

- a) an accompanying invoice or airway bill; or
- b) the physical labelling of the goods; or
- c) an overseas supplier's declaration describing the goods.

3. If the product description on the Import Permit varies from the identifying documentation provided for clearance, the importer is responsible for providing evidence to the quarantine officer that the Import Permit covers the products in the consignment.

4. The consignment must be accompanied by a declaration from the supplier or manufacturer stating that the specimens were only taken from persons with no signs or symptoms of a human quarantinable or exotic disease.

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 an AQIS 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.

POST ENTRY / END USE 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 AQIS 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 AQIS. This also applies if the product is to be used in vaccine or veterinary therapeutic manufacture.

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

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| | <p>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.</p> <p>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.</p> |

End of Condition Text