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AGREEMENT

between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products

(OJ L 57, 26.2.1997, p. 5)

Amended by:

		Official Journal		
		No	page	date
►**M1**	EXCHANGE OF LETTERS concerning the amendment to the Annexes to the Agreement between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products	L 332	3	23.12.1999
►**M2**	Agreement in the form of an Exchange of Letters concerning amendments to the Annexes to the Agreement between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products	L 333	15	10.12.2002
►**M3**	Agreement in the form of an Exchange of Letters on the amendments to the Annexes to the agreement between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products	L 214	38	26.8.2003
►**M4**	Exchange of Letters constituting an Arrangement with New Zealand on the modification of Annex V to the Agreement between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products	L 332	17	6.11.2004
►**M5**	Agreement in the form of an Exchange of Letters constituting an Arrangement with New Zealand on the amendments to Annex V and Annex VIII to the Agreement between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products	L 338	3	5.12.2006
►**M6**	Commission Implementing Decision (EU) 2015/1084 of 18 February 2015	L 175	45	4.7.2015

Corrected by:

►**C1**	Corrigendum, OJ L 313, 28.11.2003, p. 82 (22003A0826(01))

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AGREEMENT

between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products

THE EUROPEAN COMMUNITY,

of the one part, and

NEW ZEALAND,

of the other part,

hereinafter referred to as ‘the Parties’;

WHEREAS the Parties acknowledge that their systems of sanitary measures are intended to provide comparable health assurances;

REAFFIRMING their commitment to the rights and obligations established pursuant to the WTO Agreement on the Application of Sanitary and Phytosanitary Measures (hereinafter referred to as ‘the SPS Agreement’);

DESIRING to facilitate trade in live animals and animal products between the European Community (hereinafter referred to as ‘the Community’) and New Zealand while safeguarding public and animal health and thereby meeting consumer expectations in relation to the wholesomeness of food products;

DESIRING to resolve other veterinary issues applicable to trade in live animals and animal products between the Community and New Zealand;

RESOLVED to take the fullest account of the risk of spread of animal infection and disease and the measures put in place to control and eradicate such infections and diseases, and in particular to avoid disruptions to trade,

HAVE AGREED AS FOLLOWS:

Article 1

Objective

The objective of this Agreement is to facilitate trade in live animals and animal products between the Community and New Zealand by establishing a mechanism for the recognition of equivalence of sanitary measures maintained by the two Parties consistent with the protection of public and animal health, and to improve communication and cooperation on sanitary measures.

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Article 2

General provisions

The provisions set out in this Agreement shall apply in respect of trade between the Community and New Zealand in live animals and animal products.

The jointly determined arrangements for the application of this Agreement by the Parties are set out in the Annexes.

Article 3

Multilateral obligations

Nothing in this Agreement or the Annexes shall limit the rights or obligations of the Parties pursuant to the Agreement establishing the World Trade Organization and its Annexes, and in particular the SPS Agreement.

Article 4

Scope

1. The scope of this Agreement shall be limited initially to the sanitary measures applied by either Party to the live animals and animal products listed in Annex I, except as provided for in paragraphs 2 and 3.

2. Unless otherwise specified under the provisions set out in the Annexes to this Agreement and without prejudice to Article 11, this Agreement shall not apply to sanitary measures related to food additives (all food additives and colours), sanitary stamps, processing aids, flavours, irradiation (ionization), contaminants (including microbiological standards), transport, chemicals originating from the migration of substances from packaging materials, labelling of foodstuffs, nutritional labelling, medicated feeds and premixes.

3. The Parties may also agree to apply the principles of this Agreement to address veterinary issues other than sanitary measures applicable to trade in live animals and animal products.

4. The Parties may agree to modify this Agreement in the future, to extend the scope to other sanitary or phytosanitary measures affecting trade between the Parties.

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Article 5

Definitions

For the purposes of this Agreement the following definitions shall apply:

(a) live animals and animal products: means the live animals and animal products covered by the provisions listed in Annex I;

(b) sanitary measures: means sanitary measures as defined in Annex A, paragraph 1, of the SPS Agreement falling within the scope of this Agreement;

(c) appropriate level of sanitary protection: means the level of protection as defined in Annex A, paragraph 5, of the SPS Agreement;

(d) region: means ‘zones’ and ‘regions’ as defined in the Animal Health Code of the Office international des epizooties;

(e) responsible authorities:

(i) New Zealand — the authorities described in Part A of Annex II;

(ii) European Community — the authorities described in Part B of Annex II.

Article 6

Adaptation to regional conditions

1. The Parties recognize for trade between them regional freedom from the animal diseases specified in Annex III.

2. Where one of the Parties considers that it has a special status with respect to a specific disease, it may request recognition of this status. The Party concerned may also request additional guarantees in respect of imports of live animals and animal products appropriate to the agreed status. The guarantees for specific diseases shall be specified in Annex V.

3. Without prejudice to paragraph 2, the importing Party shall recognize regionalization decisions taken in accordance with criteria as defined in Annex IV as the basis for trade from a Party within which an area is affected by one or more of the diseases listed in Annex III.

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Article 7

Equivalence

1. The recognition of equivalence requires an assessment and acceptance of:

— the legislation, standards and procedures, as well as the programmes in place to allow control and to ensure domestic and importing countries' requirements are met,

— the documented structure of the relevant responsible authority(ies), their powers, their chain of command, their modus operandi and the resources available to them,

— the performance of the relevant responsible authority in relation to the control programme and assurances.

In this assessment, the Parties shall take account of experience already acquired.

2. Equivalence shall be applied in relation to sanitary measures for live animal or animal product sectors, or parts of sectors, in relation to legislation, inspection and control systems, parts of systems, or in relation to specific legislation, inspection and/or hygiene requirements.

Article 8

Determination of equivalence

1. In reaching a determination of whether a sanitary measure applied by an exporting Party achieves the importing Party's appropriate level of sanitary protection, the Parties shall follow a process that includes the following steps:

(i) the identification of the sanitary measure(s) for which recognition of equivalence is sought;

(ii) the explanation by the importing Party of the objective of its sanitary measure(s), including an assessment, as appropriate to the circumstances, of the risk, or risks, that the sanitary measure(s) is intended to address, and identification by the importing Party of its appropriate level of sanitary protection;

(iii) the demonstration by the exporting Party that its sanitary measure(s) achieves the importing Party's appropriate level of sanitary protection;

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(iv) the determination by the importing Party of whether the exporting Party's sanitary measure(s) achieves its appropriate level of sanitary protection;

(v) the importing Party shall accept the sanitary measure(s) of the exporting Party as equivalent if the exporting Party objectively demonstrates that its measure(s) achieves the importing Party's appropriate level of protection.

2. Where equivalence has not been recognized, trade may take place under the conditions required by the importing Party to meet its appropriate level of protection as set out in Annex V. The exporting Party may agree to meet the importing Party's conditions, without prejudice to the result of the process set out in paragraph 1.

Article 9

Recognition of sanitary measures

1. Annex V lists those sectors, or parts of sectors, for which, at the date of entry into force of this Agreement, the respective sanitary measures are recognized as equivalent for trade purposes. The Parties shall take the necessary legislative/administrative actions to implement recognition of equivalence to allow trade on that basis within three months.

2. Annex V also lists those sectors, or parts of sectors, for which the Parties apply differing sanitary measures and have not concluded the assessment provided for in Article 7. Based on the process described in Articles 7 and 8, the actions set out in Annex V shall be taken to enable the assessment to be completed by the indicative dates indicated therein. The Parties shall take the necessary legislative/administrative actions to implement recognition of equivalence within three months of the date of recognition. Pending recognition, trade shall take place under the conditions set out in Annex V.

3. Each consignment of live animals or animal products for which equivalence has been recognized presented for import will be accompanied, unless not required, by an official health certificate, the model attestation for which is prescribed in Annex VII. The Parties may jointly determine principles or guidelines for certification. Any such principles shall be included in Annex VII.

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Article 10

Verification

1. To maintain confidence in the effective implementation of the provisions of this Agreement, each Party shall have the right to carry out audit and verification procedures of the exporting Party, which may include:

(a) an assessment of all or part of the responsible authorities' total control programme, including, where appropriate, reviews of the inspection and audit programmes; and

(b) on-the-spot checks.

These procedures shall be carried out in accordance with the provisions of Annex VI.

2. Each Party shall also have the right to carry out frontier checks on consignments on importation, the results of which form part of the verification process.

3. For the Community:

— the Community shall carry out the audit and verification procedures provided for in paragraph 1,

— the Member States shall carry out the frontier checks provided for in paragraph 2.

4. For New Zealand, the New Zealand authorities shall carry out the audit and verification procedures and frontier checks provided for in paragraphs 1 and 2.

5. On the mutual consent of the Parties to this Agreement, either Party may:

(a) share the results and conclusions of its audit and verification procedures and frontier checks with countries that are not parties to this Agreement, or

(b) use the results and conclusions of the audit and verification procedures and frontier checks of countries that are not parties to this Agreement.

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Article 11

Frontier checks and inspection fees

1. The frequencies of frontier checks, as referred to in Article 10 (2), on imported live animals and animal products shall be as set out in Annex VIII A. The Parties may amend the frequencies, within their responsibilities, as appropriate as a result of progress made in accordance with Annex V and Annex IX, or as a result of other actions or consultations provided for in this Agreement.

2. The physical checks applied shall be based on the risk associated with such importations.

3. In the event that the checks reveal non-conformity with the relevant standards and/or requirements, the action taken by the importing Party should be based on an assessment of the risk involved. Wherever possible, the importer or his representative shall be given access to the consignment and the opportunity to contribute any relevant information to assist the importing Party in taking a final decision.

4. Inspection fees may be collected for the costs incurred in frontier checks. Provisions in relation to inspection fees are prescribed in Annex VIII B.

Article 12

Notification

1. The Parties shall notify each other of:

— significant changes in health status such as the presence and evolution of diseases in Annex III within 24 hours,

— findings of epidemiological importance with respect to diseases which are not in Annex III or new diseases without delay,

— any additional measures beyond the basic requirements of their respective sanitary measures taken to control or eradicate animal disease or protect public health, and any changes in preventive policies, including vaccination policies.

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2. The notifications referred to in paragraph 1 shall be made in writing to the contact points established in accordance with Article 15 (4).

3. In cases of serious and immediate concern with respect to public/animal health, oral notification shall be made to the contact points established in accordance with Article 15 (4), and written confirmation should follow within 24 hours.

4. Where either Party has serious concerns regarding a risk to animal or public health, consultations regarding the situation shall, on request, take place as soon as possible, and in any case within 14 days. Each Party shall endeavour in such situations to provide all the information necessary to avoid a disruption in trade, and to reach a mutually acceptable solution.

Article 13

Safeguard clause

Without prejudice to Article 12, and in particular paragraph 4, either Party may, on serious public or animal health grounds, take provisional measures necessary for the protection of public or animal health. These measures shall be notified within 24 hours to the other Party and, on request, consultations regarding the situation shall be held within 14 days. The Parties shall take due account of any information provided through such consultations.

Article 14

The principles of this Agreement shall also be applied to address outstanding issues falling within its scope affecting trade between the Parties in live animals and animal products as listed in Annex IX. Modifications shall be made to this Annex and, as appropriate, the other Annexes, to take account of progress made and new issues identified.

Article 15

Information exchange and submission of scientific research and data

1. The Parties shall exchange information relevant to the implementation of this Agreement on a uniform and systematic basis, to provide assurance, engender mutual confidence and demonstrate the efficacy of the programmes controlled. Where appropriate, achievement of these objectives may be enhanced by exchanges of officials.

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2. The information exchange on changes in their respective sanitary measures, and other relevant information, shall include:

— opportunity to consider proposals for changes in regulatory standards or requirements which may affect this Agreement in advance of their finalization. Where either Party considers it necessary, proposals may be dealt with in accordance with Article 16 (3),

— briefing on current developments affecting trade in live animals and animal products,

— information on the results of the verification procedures provided for in Article 10.

3. The Parties shall provide for the submission of scientific papers or data to the relevant scientific forums to substantiate their views/claims. Such evidence shall be evaluated by the relevant scientific forums in a timely manner, and the results of that examination shall be made available to both Parties.

4. The contact points for this exchange of information are set out in Annex X.

Article 16

Joint management committee

1. A joint management committee (hereinafter referred to as 'the Committee') consisting of representatives of the Parties shall be established, which shall consider any matters relating to the Agreement and shall examine all matters which may arise in relation to its implementation. The Committee shall meet within one year of the entry into force of this Agreement, and at least annually thereafter. The Committee may also address issues out of session by correspondence.

2. The Committee shall, at least once a year, review the Annexes to this Agreement, notably in the light of progress made under the consultations provided for under this Agreement. Modifications to the Annexes will be jointly determined.

3. The Parties may agree to establish technical working groups consisting of expert-level representatives of the Parties, which shall identify and address technical and scientific issues arising from this Agreement.

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When additional expertise is needed, the Parties may also establish *ad hoc* technical or scientific working groups, whose membership need not be restricted to representatives of the Parties.

Article 17

Territorial application

The territorial application of this Agreement shall be as follows:

(a) the Community: to the territories in which the Treaty establishing the European Community is applied and under the conditions laid down in that Treaty;

(b) New Zealand: to all territorial areas of New Zealand. However this Agreement shall not apply to Tokelau.

Article 18

Final provisions

1. This Agreement shall be approved by the Parties in accordance with their respective procedures.

This Agreement shall enter into force on the first day of the month following the date on which the Parties notify each other in writing that the procedures mentioned in the preceding subparagraph have been completed.

2. Each Party shall implement the commitments and obligations arising from this Agreement in accordance with its internal procedures.

3. Either Party may, at any time, propose amendments to this Agreement. Any agreed amendments shall enter into force on the first day of the month following the date on which the Parties notify each other in writing that their respective internal procedures for the approval of amendments have been completed.

4. Either Party may denounce this Agreement by giving at least six months' notice in writing. In such an event, the Agreement shall come to an end on the expiry of the period of notice.

5. This Agreement shall be drawn up in two copies in the English language, each of these texts being equally authentic.

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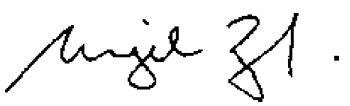
Done at Brussels, this seventeenth day of December in the year one thousand nine hundred and ninety-six.

For the European Community

For New Zealand



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LIST OF ANNEXES

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ANNEX I

LIVE ANIMALS AND ANIMAL PRODUCTS

Live animals and animal products	As defined by
1. Live cattle and pigs	Council Directive 64/432/EEC of 26 June 1964
2. Bovine semen	Council Directive 88/407/EEC of 14 June 1988,
3. Bovine embryos	Council Directive 89/556/EEC of 25 September 1989
4. Live horses	Council Directive 90/426/EEC of 26 June 1990
5. Pig semen	Council Directive 90/429/EEC of 26 June 1990
6. Poultry and hatching eggs	Council Directive 90/539/EEC of 15 October 1990
7. Live aquaculture animals and aquaculture products	Council Directive 91/67/EEC of 28 January 1991
8. Live sheep and goats	Council Directive 91/68/EEC of 28 January 1991
9. Other live animals, semen, ova and embryos from the animal species not referred to in points 1 to 8	Council Directive 92/65/EEC of 13 July 1992
10. Fresh meat	Council Directive 64/433/EEC of 26 June 1964
11. Fresh poultry meat	Council Directive 71/118/EEC of 15 February 1971
12. Meat products	Council Directive 77/99/EEC of 21 December 1976
13. Minced meat and meat preparations	Council Directive 94/65/EC of 14 December 1994
14. Egg products	Council Directive 89/437/EEC of 20 June 1989
15. Live bivalve molluscs	Council Directive 91/492/EEC of 15 July 1991
16. Fisheries products	Council Directive 91/493/EEC of 22 July 1991
17. Farmed game meat	Council Directive 91/495/EEC of 27 November 1991
18. Wild game meat	Council Directive 92/45/EEC of 16 June 1992
19. Milk and milk products	Council Directive 92/46/EEC of 16 June 1992
20. Animal waste	Council Directive 90/667/EEC of 27 November 1990
21. Animal products not referred to in points 10 to 20	Council Directive 92/118/EEC of 17 December 1992

Note:

Under New Zealand legislation (Biosecurity Act (1993) and the ‘saved provisions’ of the Animals Act 1967) a list is prescribed of organisms that are prohibited entry into New Zealand.

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ANNEX II

RESPONSIBLE AUTHORITIES

PART A

New Zealand

The Ministry for Primary Industries is responsible for controls in sanitary issues and veterinary affairs.

— In terms of exports to the European Union the Ministry for Primary Industries is responsible for setting sanitary (food safety) and animal health (zoosanitary) standards and requirements and specifying the health certification attesting to the agreed sanitary and zoosanitary standards and requirements,

— In terms of imports into New Zealand, the Ministry for Primary Industries is responsible for setting sanitary (food safety) and animal health (zoosanitary) standards and requirements.

PART B

European Union

Control is shared between the national services in the individual Member States and the European Commission. In this respect, the following applies:

— In terms of exports to New Zealand, the Member States are responsible for the control of production circumstances and requirements, including statutory inspections/audits and issuing health certification attesting to the agreed standards and requirements,

— In terms of imports into the European Union, the European Commission is responsible for overall coordination, inspections/audits of control systems and the necessary legislative action to ensure uniform application of standards and requirements within the internal market.

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ANNEX III

DISEASES FOR WHICH REGIONALIZATION DECISIONS CAN BE TAKEN

LEGAL BASIS

Disease	EC	NZ
Foot-and-mouth disease	85/511, 64/432	Biosecurity Act parts IV, V, VI, VII and VIII
Swine vesicular disease	92/119, 64/432	Biosecurity Act parts IV, V, VI, VII and VIII
Vesicular stomatitis	92/119	Biosecurity Act parts IV, V, VI, VII and VIII
African horse sickness	90/426, 92/35	Biosecurity Act parts IV, V, VI, VII and VIII
African swine fever	64/432	Biosecurity Act parts IV, V, VI, VII and VIII
Bluetongue	92/119	Biosecurity Act parts IV, V, VI, VII and VIII
Highly pathogenic Avian influenza	92/40, 90/539	Biosecurity Act parts IV, V, VI, VII and VIII
Newcastle disease	92/66, 90/539	Biosecurity Act parts IV, V, VI, VII and VIII
Peste des petits ruminants	92/119	Biosecurity Act parts IV, V, VI, VII and VIII
Rinderpest	92/119, 64/432	Biosecurity Act parts IV, V, VI, VII and VIII
Classical swine fever	80/217, 64/432	Biosecurity Act parts IV, V, VI, VII and VIII
Contagious bovine pleuropneumonia	64/432	Biosecurity Act parts IV, V, VI, VII and VIII
Sheep pox	92/119	Biosecurity Act parts IV, V, VI, VII and VIII
Rift Valley fever	92/119	Biosecurity Act parts IV, V, VI, VII and VIII
Lumpy skin disease	92/119	Biosecurity Act parts IV, V, VI, VII and VIII

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Disease	EC	NZ
Infectious haematopoietic necrosis (IHN) (*)	91/67	Biosecurity Act parts IV, V, VI, VII and VIII
Spring viraemia of carp (SVC) (*)	91/67	Biosecurity Act parts IV, V, VI, VII and VIII
Viral haemorrhagic septicaemia (VHS) (*)	91/67	Biosecurity Act parts IV, V, VI, VII and VIII

(*) New Zealand has concerns about the ability to regionalize these diseases and will assess information and perform a risk assessment as to the technical basis for recognition by December ►M1 1999 ◄.

Asterisks for SVD, ND, AI, CSF, have been removed by New Zealand although special trade conditions for these diseases may remain in the interim — refer to Annex V for specific details.

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ANNEX IV

REGIONALIZATION AND ZONING

The Parties have jointly determined that the following forms the basis for regionalization decisions for the diseases listed in accordance with Annex III. Each Party will recognize regionalization decisions taken in accordance with the standard contained within this Annex.

In assessing risk from a given proposed importation of animals or animal products, three sets of factors may be considered:

1. Source risk factors

2. Commodity risk factors

3. Destination risk factors

Source risk factors

The primary determinant of the risk of importing disease is the status of the country of origin in respect of the disease in question. However, declarations of disease freedom must be backed up by effective surveillance programmes.

The overriding consideration in this context, therefore, is the quality of the veterinary infrastructure. No other factors can be assessed without full confidence in the veterinary administration. In particular, their ability to detect and control an outbreak of disease and to provide meaningful certification is crucial.

The ability to detect the presence of disease depends on the surveillance carried out. This surveillance can be active, passive, or both.

Active surveillance implies definitive action intended to identify the presence of disease, such as systematic clinical inspections, *ante*- and *post-mortem* examination, serology on farm or in abattoir, referral of pathological material for laboratory diagnosis, sentinel animals.

Passive surveillance means that the disease must be compulsorily notifiable, and that there must be a sufficiently high level of supervision or the animals in order to ensure that the disease will be observed quickly and reported as a suspect. There must also be a mechanism for investigation and confirmation, and a high level of awareness of the disease and its symptoms by farmers and vets.

Epidemio-surveillance may be augmented by voluntary and compulsory herd/flock health programmes, particularly those which ensure a regular veterinary presence on the farm.

Other factors to be considered include:

— disease history,

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— vaccination history,

— controls on movements into the zone, out of the zone and within the zone,

— animal identification and recording,

— presence of disease in adjacent areas,

— physical barriers between zones of differing status,

— meteorological conditions,

— use of buffer zones (with or without vaccination),

— presence of vectors and/or reservoirs,

— active control and eradication programmes (where appropriate),

— *ante- and post-mortem* inspection system.

On the basis of these factors, a zone may be defined.

The authority with the responsibility for implementing the zoning-policy is in the best position to define and maintain the zone. When there is a high level of confidence in that authority, the decisions it makes can be the basis for trade.

The zones so defined may be assigned a risk category.

Possible categories are:

— low/negligible risk,

— medium risk,

— high risk,

— unknown risk.

Calculation of estimates of risk for, for example live animals may assist in this categorization. Import conditions may then be defined for each category, disease and commodity, individually or in groups.

Low/negligible risk implies that importation may take place based on a simple guarantee of origin.

Medium risk implies that some combination of certification and/or guarantees may be required before or after importation.

High risk implies that importation will only take place under conditions which significantly reduce the risk, for example by additional guarantees, testing or treatment.

Unknown risk implies that imports will only take place if the commodity itself is of very low risk, for example hides, wool, or under the conditions for ‘high risk’ if the commodity factors warrant.

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Commodity risk factors

These include:

— is the disease transmissible by the commodity?

— could the agent be present in the commodity if derived from a healthy and/or clinically affected animal?

— can the predisposing factor be reduced, for example by vaccination?

— what is the likelihood that the commodity has been exposed to infection?

— has the commodity been obtained in such a way as to reduce the risk, for example deboning?

— has the commodity been subjected to a treatment which inactivates the agent?

Appropriate tests and quarantine will reduce the risk.

Destination risk factors

— presence of susceptible animals,

— presence of vectors,

— possible vector-free period,

— preventive measures such as waste food feeding and animal waste rendering rules,

— intended use of product, for example petfood, human consumption only.

These factors are inherent in or are under the control of the importing country, and some may therefore be modified to facilitate trade. These may for example include restricted entry conditions, for example animals to be confined to a certain vector-free region until the incubation period has passed, or canalization systems.

However, destination risk factors will also be taken into account by the infected country with respect to the risk presented by movements from the infected part to the free part of its territory.

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ANNEX V

RECOGNITION OF SANITARY MEASURES

Glossary

Yes (1)	Equivalence agreed. Model health attestations in Annex VII, Section 1(a) to be used. The EU may lay down its import certificates for live animals and animal products from New Zealand with a 'Yes-1' status in TRACES using a model as agreed by both Parties.
Yes (2)	Equivalence agreed in principle. Some specific issue(s) to be resolved. Importing party's model health certificate or veterinary documents to be used.
Yes (3)	Equivalence in form of compliance with importing Party's requirements. Importing party's model health certificate or veterinary documents to be used.
NE	Not evaluated. Importing party's model veterinary health certificate or veterinary documents to be used.
E	Still evaluating — under consideration. Importing party's model health certificate or veterinary documents to be used.
[]	Issues targeted for imminent resolution.
No	Not equivalent and/or further evaluation is required. Trade may occur if the exporting Party meets the importing Party's requirements.
N.A.	Not applicable
ASF	African swine fever
BSE	Bovine spongiform encephalopathy
BT	Bluetongue
C	Celsius
CBPP	Contagious bovine pleuropneumonia
CSF	Classical swine fever
EU/NZ	European Union/New Zealand
E-Cert	New Zealand's electronic data transmission system for export health certification.
EIA	Equine infectious anaemia
FMD	Foot and mouth disease
gst	goods and services tax
HPNAI	Highly pathogenic notifiable avian influenza
HTST	High Temperature/Short Time

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IBR	Infectious bovine rhinotracheitis
LPNAI	Low pathogenic notifiable avian influenza
LSD	Lumpy skin disease
min	minute(s)
ND	Newcastle disease
None	No special conditions
OIE	Office International des Epizooties
PAP	Processed animal protein
PPR	Peste des petits ruminants
PRRS	Porcine Reproductive and Respiratory Syndrome
RND	Rinderpest
SVD	Swine vesicular disease
TRACES	The EU's electronic data transmission system for (export) health certification.
TSE	Transmissible spongiform encephalopathy
UHT	Ultra high temperature
VS	Vesicular stomatitis

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Section 1

Germplasm and live animals

Commodity	EU Exports to New Zealand (1)					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
1. **Semen**										
— Cattle	88/407/EEC	Biosecurity Act 1993 S 22	Yes (1)	See Chapter 28: — Q-fever — Blue-tongue		Animal Products Act 1999	88/407/EEC 2011/630/EU	E	IBR. see Chapter 28	The EU to consider reviewing whether testing of semen for IBR using the OIE approved PCR testing methodology provides an equivalent assurance to IBR disease freedom.
— Sheep/goats	92/65/EEC 2010/470/EU	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	92/65/EEC 2010/472/EU	NE		
— Pigs	90/429/EEC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	90/429/EEC 2012/137/EU	NE		

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Commodity	EU Exports to New Zealand [1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Deer	92/65/EEC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	92/65/EEC	No		
— Horses	92/65/EEC 2010/470/EU	Biosecurity Act 1993 S 22	Yes (3)			Animal Products Act 1999	92/65/EEC 2004/211/EC 2010/471/EU	Yes (3)		
— Dogs	92/65/EEC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	92/65/EEC	NE		

2. **Embryos (except embryos subject to penetration of the *zona pellucida*)**

— Cattle *in-vivo* derived embryos	89/556/EEC	Biosecurity Act 1993	Yes (1)	See Chapter 28: — Q-fever — Bovine viral diarrhoea (type II)		Animal Products Act 1999	89/556/EEC 2006/168/EC	Yes (1)		
in vitro derived embryos	89/556/EEC	Biosecurity Act 1993	Yes (1)	See Chapter 28: — Q-fever — Bovine viral diarrhoea (type II)		Animal Products Act 1999		Yes (3)		

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Commodity	EU Exports to New Zealand [1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Sheep/goats	92/65/EEC 2010/470/EU	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	92/65/EEC 2010/472/EU	NE		
— Pigs	92/65/EEC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	92/65/EEC	NE		
— Deer	92/65/EEC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	92/65/EEC	No		
— Horses	92/65/EEC 2010/470/EU	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	92/65/EEC 2004/211/EC 2010/471/EU	Yes (3)		
— Poultry hatching eggs	2009/158/EC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2009/158/EC Regulation (EC) No 798/2008	Yes (3)	Salmonella see Chapter 28.	
— Ratites hatching eggs								NE		
3. **Live animals**										
— Cattle	64/432/EEC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	Regulations (EC) No 999/2001 (EU) No 206/2010	Yes (3)	IBR see Chapter 28	

▼M6

Commodity	EU Exports to New Zealand [1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Sheep/goats	91/68/EEC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2004/212/EC Regulations (EC) No 999/2001 (EU) No 206/2010	Yes (3)		The EU to consider scrapie freedom of NZ
— Swine	64/432/EEC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulation (EU) No 206/2010	Yes (3)	Aujeszky's disease see Chapter 28	
— Deer	2004/68/EC 92/65/EEC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	2004/68/EC Regulation (EU) No 206/2010	Yes (3)		
— *Equidae*	2009/156/EC	Biosecurity Act 1993 S 22	Yes (3)			Animal Products Act 1999	92/260/EEC 93/195/EEC 93/196/EEC 93/197/EEC 2004/211/EC 2009/156/EC 2010/57/EU	Yes (3)	EIA see Chapter 28	

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Commodity	EU Exports to New Zealand [1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Dogs, cats and ferrets	Commercial: 92/65/EEC 2013/519/EU Non-commercial: 2003/803/EC Regulations (EC) No 998/2003 (EU) No 576/2013 (EU) No 577/2013	Biosecurity Act 1993 S 22	Yes (3)	Rabies see Chapter 28		Animal Products Act 1999	Commercial Imports: 92/65/EEC 2011/874/EU 2013/519/EU Non-commercial: 2011/874/EU 2013/519/EU 2013/520/EU Regulations (EC) No 998/2003 (EU) No 576/2013 (EU) No 577/2013	Yes (3)	Rabies see Chapter 28	
— Live poultry	2009/158/EC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2009/159/EC Regulation (EC) No 798/2008	Yes (3)	Salmonella see Chapter 28	

▼M6

Commodity	EU Exports to New Zealand (1)					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Ratites			NE					NE		
— Live bees bumble bees including bee/bumble bee germplasm	92/65/EEC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	92/65/EEC 2013/503/EU Regulation (EU) No 206/2010	Yes (1)	Bees/bumble bees see Chapter 28 No trade of commodities to Member States or regions, listed in the Annex to Commission Implementing Decision 2013/503/EU.	

(1) Commodities must be fully eligible for unrestricted intra-Union trade, unless otherwise indicated.

▼M6

Section 2

Meat (including fresh meat, fresh poultry meat, farmed and wild game meat), minced meat, meat preparations and meat products for human consumption

Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
4. **Meat**										
4.A. **Fresh Meat as defined in Regulation (EC) No 853/2004.** Includes minced meat unprocessed (fresh) blood/bones/fat for human consumption.										
Animal health — Ruminants — Equidae	2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	2002/99/EC Regulations (EC) No 999/2001 (EU) No 206/2010	Yes (1)		
— Pigs	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	2002/99/EC Regulation (EU) No 206/2010	Yes (1)		
Public health	Regulations (EC) (1) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	2011/163/EU Regulations (EC) (1) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EU) No 206/2010	Yes (1)	Salmonella and BSE see Chapter 28 — Minced meat must be frozen.	

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Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
4.B. **Fresh Poultry Meat**										
Animal health — poultry	2002/99/EC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2002/99/EC Regulation (EC) No 798/2008	Yes (3)		
— turkeys			Yes (3)					NE		
Public health	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	NE		
4.C. **Farmed Game Meat**										
Animal health — Deer — Pigs	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	2002/99/EC Regulation (EU) No 206/2010	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Rabbit	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	2002/99/EC Regulation (EC) No 119/2009	Yes (1)		
— Other land mammals	2002/99/EC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	2002/99/EC Regulation (EC) No 119/2009	Yes (1)		
— Feathered (including ratite)	2002/99/EC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2002/99/EC Regulation (EC) No 798/2008	Yes (3)		
Public health — Land mammals	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 119/2009	Yes (1)		

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Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU Regulation (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	Yes (3)		
— Ratite			Yes (1)					Yes (1)		

4.D. **Wild game meat**

Animal health — Deer — Rabbit	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	2002/99/EC Regulations (EC) No 119/2009 (EU) No 206/2010	Yes (1)		
— Pigs	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	CSF and PRRS see Chapter 28		Animal Products Act 1999	2002/99/EC Regulations (EC) No 119/2009 (EU) No 206/2010	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Other wild land mammals	2002/99/EC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	2002/99/EC Regulation (EC) No 119/2009	NE		
— Feathered	2002/99/EC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	2002/99/EC Regulation (EC) 798/2008	Yes (3)		
Public health — Wild land mammals	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 119/2009	Yes (1)	Unskinned and uneviscerated wild *leporidae* must be chilled to + 4 °C for a maximum of 15 days prior to the intended time of import.	

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Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	NE		

5. **Meat preparations**

5.A. **Meat preparations from fresh meat**

Animal health — Ruminants — Pigs	2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	2000/572/EC 2002/99/EC Regulation (EC) No 999/2001	Yes (1)		
Public health	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	2000/572/EC 2011/163/EU Regulation (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)	Frozen only BSE see Chapter 28	

▼M6

Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
5.B. **Meat preparations derived from fresh poultry meat**										
Animal health — Poultry	2002/99/EC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2000/572/EC 2002/99/EC Regulation (EC) No 798/2008	Yes (3)		
— Turkey			Yes (3)					NE		
Public health	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956 Animal Products Act 1999	Yes (1)			Animal Products Act 1999	2000/572/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	NE	Frozen only	
5.C. **Meat preparations derived from farmed game meat**										
Animal health — Deer — Pigs	92/118/EEC 2002/99/EC Regulation (EU) No 206/2010	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2000/572/EC 2002/99/EC Regulation (EU) No 206/2010	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Rabbit	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	92/118/EEC 2000/572/EC 2002/99/EC	Yes (1)		
— Ratites	2002/99/EC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2000/572/EC 2002/99/EC Regulation (EC) No 798/2008	Yes (3)		
— Feathered	2002/99/EC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2000/572/EC 2002/99/EC Regulation (EC) No 798/2008	Yes (3)		
Public health — Deer — Pigs — Rabbit	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2000/572/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)	Frozen only	

▼M6

Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered — Ratites	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004		Yes (1)				2000/572/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	NE Yes (1)		

5.D. **Meat preparations derived from wild game meat**

Animal health — Deer — Rabbit	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	2000/572/EC 2002/99/EC	Yes (1)		
— Pigs	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	CSF and PRRS see Chapter 28		Animal Products Act 1999	2000/572/EC 2002/99/EC	Yes (1)		
— Feathered	2002/99/EC	Biosecurity Act 1993 S 22	No			Animal Products Act 1999	2000/572/EC 2002/99/EC Regulation (EC) No 798/2008	Yes (3)		

▼M6

Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health — Wild land mammals	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2000/572/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)	Frozen Only	
— Feathered	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004		Yes (1)				2000/572/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	NE		

6. **Meat products**

6.A. **Meat products derived from fresh meat**

Animal health — Ruminants — Horses — Pigs	2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	2002/99/EC 2007/777/EC Regulation (EC) No 999/2001	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)	BSE see Chapter 28	

6.B. **Meat products derived from fresh poultry meat**

Animal health	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0 3$ treatment		Animal Products Act 1999	2002/99/EC 2007/777/EC Regulation (EC) No 798/2008	Yes (3)		

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Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	NE		

6.C. **Meat products derived from farmed game**

Animal health — Pigs — Deer — Rabbit	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC	Yes (1)		
— Ratites	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0 3$ treatment		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 798/2008	Yes (3)		

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Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Other feathered	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0$3 treatment		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC	Yes (3)		
Public health — Pigs — Deer — Rabbit	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 999/2001	Yes (1)		
— Feathered	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	Yes (3)		

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Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Ratite			Yes (1)					Yes (1)		

6.D. **Meat products derived from wild game**

Animal health Wild game — Pigs	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	CSF and PRRS see Chapter 28		Animal Products Act 1999	2002/99/EC 2007/777/EC	Yes (1)		
— Deer — Rabbit	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	2002/99/EC 2007/777/EC	Yes (1)		
— Feathered	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0 3$ treatment		Animal Products Act 1999	2002/99/EC 2007/777/EC Regulation (EC) 798/2008	Yes (3)		
Public health Wild game	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		

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Commodity	EU Exports to New Zealand					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	NE		

(1) All entries referring to Regulations (EC) No 852/2004, (EC) No 853/2004, and (EC) No 854/2004 shall be construed to include relevant implementing measures and microbial criteria as laid down in Regulations (EC) No 2073/2005, (EC) No 2074/2005 and (EC) No 2076/2005.

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Section 3

Other products for human consumption

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
7. **Products intended for human consumption**										
7.A. **Animal casings**										
Animal health — Cattle — Sheep — Goats — Pigs	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	92/118/EEC 2003/779/EC 2007/777/EC 477/2010/EU Regulation (EC) No 999/2001	Yes (1)		
Public health	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)	BSE see Chapter 28	

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
7.B. **Processed bones and bone products for human consumption**										
Animal health Fresh meat: — Ruminants — Horses — Pigs	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 999/2001	Yes (1)		
— Poultry	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0$3 treatment		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC	Yes (3)		
Farmed game — Pigs — Deer	92/118/EC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC	Yes (1)		
— Feathered	92/118//EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0$3 treatment		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC	Yes (3)		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Wild game — Deer — Pigs	92/118/EC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	CSF and PRRS see Chapter 28		Animal Products Act 1999	92/118/EC 2002/99/EC 2007/777/EC	Yes (1)		
— Feathered			Yes (1)	Heat treated shelf stable $F_0$3 treatment				Yes (3)		
Public health Fresh meat: — Ruminants — Horses — Pigs	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	2007/777/EC Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)	BSE see Chapter 28	

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Poultry Fresh meat	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999 Food Act 1981 Health Act 1956	2007/777/EC Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	NE		
Farmed game — Mammals	92/118/EEC Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999 Food Act 1981 Health Act 1956	2007/777/EC Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		
— Feathered			Yes (1)					NE		
Wild game — Mammals	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999 Food Act 1981 Health Act 1956	2007/777/EC Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered			Yes (1)					NE		

7.C. **Processed animal protein for human consumption**

Animal health PAP derived from fresh meat: — Ruminants — Horses — Pigs	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC 477/2010/EU Regulation (EC) No 999/2001	Yes (1)		
Poultry PAP derived from fresh meat	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0$3 treatment		Animal Products Act 1999	94/438/EC 92/118/EEC 2002/99/EC 2007/777/EC	Yes (3)		
Farmed game — Pigs — Deer	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC 477/2010/EU Regulation (EC) No 999/2001	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered			Yes (1)	Heat treated shelf stable $F_0$3 treatment				Yes (3)		
Wild game — Pigs — Deer	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	CSF and PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC 477/2010/EU Regulation (EC) No 999/2001	Yes (1)		
— Feathered			Yes (1)	Heat treated shelf stable $F_0$3 treatment				Yes (3)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health PAP derived from fresh meat — Ruminants — Horses — Pigs	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 999/2001	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 999/2001	Yes (1)	BSE see Chapter 28	
Poultry PAP derived from fresh meat	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	NE		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Farmed game	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	92/118/EEC 2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		
— Feathered			Yes (1)					NE		
Wild game	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered			Yes (1)					NE		

7.D. **Blood and blood products for human consumption**

Animal health Blood and blood products derived from fresh meat: — Ruminants — Horses — Pigs	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulations (EC) No 999/2001 (EU) No 206/2010	Yes (1)		
Poultry Blood and blood products fresh poultry meat	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0$3 treatment		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 798/2008	Yes (3)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Farmed game — Pigs — Deer	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 999/2001	Yes (1)		
— Feathered			Yes (1)	Heat treated shelf stable $F_0$3 treatment				Yes (3)		
Wild game — Pigs — Deer	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	CSF and PRRS see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 999/2001	Yes (1)		
— Feathered			Yes (1)	Heat treated shelf stable $F_0$3 treatment				Yes (3)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health — Ruminants — Horses — Pigs Fresh meat	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)	BSE see Chapter 28	
Poultry Fresh meat	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	NE		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Farmed game — Mammals	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		
— Feathered game			Yes (1)					NE		
Wild game — Mammals	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered game			Yes (1)					NE		

7 E. **Lard and rendered fats for human consumption**

Animal health Domestic mammals Products derived from fresh meat: — Ruminants — Horses — Pigs	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 999/2001	Yes (1)		
Poultry Products derived from fresh meat:	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)	Heat treated shelf stable $F_0 3$ treatment		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 798/2008	Yes (3)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Farmed game — Pigs — Deer	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 999/2001	Yes (1)		
— Feathered game			Yes (1)	Heat treated shelf stable $F_0$3 treatment				Yes (3)		
Wild game — Pigs — Deer	92/118/EEC 2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (1)	CSF see Chapter 28		Animal Products Act 1999	92/118/EEC 2002/99/EC 2007/777/EC Regulation (EC) No 999/2001	Yes (1)		
— Feathered game			Yes (1)	Heat treated shelf stable $F_0$3 treatment				Yes (3)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health — Ruminants — Horses — Pigs Fresh meat	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)	BSE see Chapter 28	
Poultry Fresh meat	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	NE		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Farmed game	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		
— Feathered game			Yes (1)					NE		
Wild game	92/118/EEC Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	92/118/EEC 2007/777/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Feathered game			Yes (1)					NE		

7.F. **Gelatines for human consumption**

Animal health	2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	2002/99/EC Regulation (EC) No 999/2001	NE		
Public health	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	NE	BSE see Chapter 28		Animal Products Act 1999 Food Act 1981 Health Act 1956	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 2074/2005	NE	BSE see Chapter 28	

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
7.G. **Collagen for human consumption**										
Animal health	Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulation (EC) No 999/2001	NE		
Public health	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Animal Products Act 1999 Food Act 1981 Health Act 1956	NE	BSE see Chapter 28		Animal Products Act 1999	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	NE	BSE see Chapter 28	
7.H. **Stomachs and Bladders (Salted, Dried, or heated and other products)**										
Animal health — Cattle — Sheep — Goats — Pigs	2002/99/EC Regulation (EC) No 999/2001	Biosecurity Act 1993 S 22	Yes (3)			Animal Products Act 1999	2002/99/EC 2007/777/EC Regulation (EC) No 999/2001	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health	Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU Regulations (EC) No 999/2001 (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Yes (1)		

8. **Milk and milk products for human consumption.** Includes colostrum and colostrum-based products for human consumption.

Animal health Domestic mammals including — Cattle — Buffalo — Sheep — Goats	2002/99/EC	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	2002/99/EC Regulation (EU) No 605/2010	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health — Pasteurised	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EU) No 605/2010	Yes (1)		
— Not pasteurised, thermised cheeses	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 NZ (milk and milk products processing) food standards 2002	Yes (1)	Thermised cheeses see Chapter 28		Food Act 1981 Animal Products Act (1999)	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EU) No 605/2010	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Unpasteurised milk products (excluding raw milk)	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	Yes (1)			Food Act 1981 Animal Products Act (1999)	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EU) No 605/2010	Yes (1)		

9. **Fishery products for human consumption (excluding live)**

Animal Health Wild marine — Finfish — Eggs/roes — Molluscs — Echinoderms — Tunicates, gastropods and crustaceans	2002/99/EC Regulation (EC) No 1251/2008	Biosecurity Act 1993 S 22	Yes (1)	Salmonids see Chapter 28 Eggs/roes see Chapter 28		Animal Products Act 1999	Regulation (EC) No 1251/2008	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Wild fresh water — Salmonids — Eggs/roes — Crayfish	2002/99/EC Regulation (EC) No 1251/2008	Biosecurity Act 1993 S 22	Yes (1)	Salmonids see Chapter 28 Eggs/roes see Chapter 28 Crayfish (frozen or processed)		Animal Products Act 1999	Regulation (EC) No 1251/2008	Yes (1)	Crayfish (frozen or processed)	
— Finfish (non salmonid) — Molluscs — Crustaceans	2002/99/EC Regulation (EC) No 1251/2008	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulation (EC) No 1251/2008	Yes (1)		
Aquaculture products (marine & fresh water — farmed) — Salmonids — Eggs/roes	2002/99/EC Regulation (EC) No 1251/2008	Biosecurity Act 1993 S 22	Yes (1)	Salmonids see Chapter 28 Eggs/roes see Chapter 28		Animal Products Act 1999	Regulation (EC) No 1251/2008	Yes (1)	Salmonids (gutted)	

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Molluscs, echinoderms, — Tunicates, gastropods and crustaceans	2002/99/EC Regulation (EC) No 1251/2008	Biosecurity Act 1993 S 22	Yes (1)	Frozen or processed		Animal Products Act 1999	Regulation (EC) No 1251/2008	Yes (1)	Frozen or processed	
— Finfish (non salmonid)	2002/99/EC Regulation (EC) No 1251/2008	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulation (EC) No 1251/2008	Yes (1)		
Public Health — Finfish — Eggs/roes — Bivalve molluscs, echinoderms, tunicates, gastropods and crustaceans	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU (Aquaculture) Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 2074/2005	Yes (1)		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
10. **Live fish, molluscs, crustaceans, including eggs and gametes**										
Animal health For human consumption — live molluscs echino-derms, tuni-cates, gastropods — live crus-taceans — live finfish — other aquatic animals	93/53/EEC 95/70/EC 2002/99/EC Regulation (EC) No 1251/2008	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulation (EC) No 1251/2008	Yes (1)		
For breeding, farming, rearing, relaying — live molluscs and fish	93/53/EEC 95/70/EC Regulation (EC) No 1251/2008	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulation (EC) No 1251/2008	Yes (3)		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health — live finfish — live molluscs, echino-derms, tuni-cates, gastropods — live crus-taceans — other fish	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	Yes (1)			Animal Products Act 1999	2011/163/EU (aquaculture for human consumption) Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 2074/2005	Yes (1)		

11. **Miscellaneous products for human consumption**

11.A. **Honey**

Animal health	92/118/EEC 2002/99/EC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	92/118/EEC 2002/99/EC	Yes (3)		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health	2001/110/EC Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	NE			Animal Products Act 1999	2001/110/EC 2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 2074/2005	Yes (3)		

11.B. **Frogs' legs**

Animal health	2002/99/EC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	2002/99/EC	NE		
Public health	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	NE			Animal Products Act 1999	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 2074/2005	NE		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
11.C. **Snails for human consumption**										
Animal health	2002/99/EC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	2002/99/EC	NE		
Public health	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	NE			Animal Products Act 1999	Regulations (EC) No 853/2004 (EC) No 854/2004 (EC) No 2074/2005	NE		
11.D. **Egg products**										
Animal health	2002/99/EC 2009/158/EC	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	2002/99/EC 2009/158/EC	NE		
Public health	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004	Food Act 1981 Health Act 1956	NE			Animal Products Act 1999	2011/163/EU Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 798/2008	NE		

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Section 4

Products not intended for human consumption

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
12. **Animal casings for the production of pet food or for technical purposes**										
Animal health — Cattle — Sheep — Goats — Pigs	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (2)	TSE related restrictions apply.		Animal Products Act 1999	2003/779/EC Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Yes (1)	BSE see Chapter 28	
Public health	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Health Act 1956 Agricultural Compounds and Veterinary Medicines Act 1997	Yes (1)	BSE see Chapter 28			Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Yes (1)	BSE see Chapter 28	

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
13. **Milk, milk products and colostrum not intended for human consumption**										
Animal health — Cattle — Sheep — Goats Pasteurised, UHT or sterilised	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		
Unpasteurised colostrum and milk for uses outside the feed chain	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (3)			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (3)		
Public health			N.A.					N.A.		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
14. **Bones and bone products (excluding bone meal), horns and horn products (excluding horn meal) and hooves and hoof products (excluding hoof meal) for uses other than as feed material, organic fertiliser or soil improver**										
Animal health	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Yes (1)	BSE see Chapter 28	
Public health			N.A.					N.A.		
15. **Processed (rendered) animal protein for feedingstuffs**										
Animal health PAP intended for the production of petfood	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011		Yes (1)	BSE see Chapter 28		Animal Products Act 1999	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Yes (1)	BSE see Chapter 28	

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
PAP derived from non-mammalian material	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22				Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011			
— fish material			Yes (1)					Yes (1)		
— avian material			Yes (2)	70 °C/50 min 80 °C/9 min or 100 °C/1 min or equivalent				Yes (1)		
Public health			N.A.					N.A.		

16. **Processed blood and blood products (excluding serum from *equidae*) for uses outside the feed chain**

Animal health — Bovine, ovine, caprine, porcine	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)	BSE see Chapter 28	

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— *Equidae*	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE		
— Avian	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE		
Public health			N.A.					N.A.		

17. **Lard and rendered fats not for human consumption, including fish oils**

Animal health — rendered fats and oils	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)	BSE see Chapter 28 Additional BSE-related labelling requirements apply.		Animal Products Act 1999	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Yes (1)	BSE see Chapter 28	

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
— Lards (porcine)	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)	Product to be derived from porcine fresh meat, farmed and wild game with Yes (1) for animal health indicated previously.		Animal Products Act 1999	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		
				CSF see Chapter 28						
— Fish oil	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Fat derivatives from Cat 2 or Cat 3 material as in Regulation (EC) No 1069/2009	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Biosecurity Act 1993 S 22	Regulations (EC) No 1069/2009 (EU) No 142/2011	E		
Public health			N.A.					N.A.		

18. A. **Gelatines for feed or for purposes outside the feed chain**

Animal health	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE		
Public health			N.A.					N.A.		

18. B. **Hydrolysed Protein, collagen, di and tri-calcium phosphate**

Animal health	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health			N.A.					N.A.		

19. **Hides and skins**

Animal health — Ungulates excluding *equidae*	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		
— Equidae — Other mammals	Regulations (EC) No 1069/2009. (EU) No 142/2011		NE				Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		
— Ratite (Ostrich, emu, rhea)	Regulation (EC) No 1069/2009	Biosecurity Act 1993 S 22	NE				Regulation (EC) No 1069/2009	Yes (1)		

▼M6

Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health			N.A.					N.A.		

20. **Wool and fibre/hair**

Animal health — Sheep, goats, camelids	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)	Scoured wool only	Clean and washed at 75 °C or equivalent	Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		
— Other ruminants and pigs	Regulations (EC) No 1069/2009 (EU) No 142/2011		NE				Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		
— Other	Regulations (EC) No 1069/2009 (EU) No 142/2011		NE				Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health			N.A.					N.A.		

21. **Petfood (includes processed) containing only category 3 material**

Animal health Processed petfood (mammalian) Hermetically sealed containers Semi-moist and dried petfood dog chews from ungulates (excluding *equidae*)	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)	BSE see Chapter 28		Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)	BSE see Chapter 28	

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Flavouring innards			NE					Yes (3)		
Processed petfood (non-mammalian) — Hermetically sealed containers — Semi-moist and dried petfood	Regulations (EC) No 1069/2009 (EU) No 142/2011		Yes (1)				Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		
— fish material			Yes (1)					Yes (1)		
— avian material			Yes (2)	70 °C/50 min 80 °C/9 min 100 °C/1 min or equivalent				Yes (1)		
Flavouring innards			NE					Yes (3)		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Raw petfood For direct consumption	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE	BSE see Chapter 28	
Public health			N.A.					N.A.		

22. **Serum from *equidae***

Animal health	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE		
Public health			N.A.					N.A.		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
23. **Other animal by-products for the manufacture of feed including petfood, and for uses outside the feed chain**										
Animal health Fresh meat — Bovine — Ovine — Caprine — Porcine — Equine	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)	Product to be derived from fresh meat, farmed and wild game with Yes (1) for animal health indicated previously		Animal Products Act 1999	Regulations (EC) No 999/2001 (EC) No 1069/2009 (EU) No 142/2011	Yes (1)	BSE see Chapter 28	
Farmed game — Pigs — Deer Wild game — Pigs — Deer				BSE see Chapter 28 Additional BSE-related labelling requirements apply						
				CSF see Chapter 28						

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Fresh meat — Poultry	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE		
Farmed and wild game — Feathered										
Other species	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	E		
Public health			N.A.					N.A.		

24. **Apiculture products — not for human consumption**

Animal health	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE		

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Commodity	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Public health			N.A.					N.A.		

25. **Game trophies**

Animal health — Mammalian	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	Yes (1)			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	Yes (1)		
— Avian			NE					NE		
Public health			N.A.					N.A.		

26. **Manure — processed**

Animal health	Regulations (EC) No 1069/2009 (EU) No 142/2011	Biosecurity Act 1993 S 22	NE			Animal Products Act 1999	Regulations (EC) No 1069/2009 (EU) No 142/2011	NE		
Public health			N.A.					N.A.		

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Section 5

General horizontal issues

	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
27. **Horizontal issues**										
Water	98/83/EC	Animal Products Act 1999 Health Act 1956	Yes (1)			Animal Products Act 1999	98/83/EC	Yes (1)		
Residues Residue monitoring — Red meat species	96/22/EC 96/23/EC	Animal Products Act 1999 Food Act 1981	Yes (1)			Animal Products Act 1999	96/22/EC 96/23/EC	Yes (1)		
— Other species other products			Yes (3)					Yes (3)		

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	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Certification Systems	96/93/EC	Animal Products Act 1999	Yes (1) Equivalence status applies to all animals and animal product commodities accorded both animal and public health equivalence 'Yes (1)' as appropriate.			Animal Products Act 1999	92/118/EEC 96/93/EC 2002/99/EC Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 2074/2005 (EC) No 1251/2008 (EC) No 1069/2009 (EU) No 142/2011	Yes (1) Equivalence status applies to animals and animal product commodities listed with 'Yes (1)' equivalence status under entry Numbers 3, 4A, 4C, 4D, 5A, 5C, 5D, 6A, 6C, 6D, 7A, 7B, 7C, 7D, 7E, 7H, 9, 10, 12, 15, 16, 17, 19, 21 and 23	When the official health certificate is issued after the departure of the consignment, it shall include reference to the appropriate Eligibility Document (ED), date of issuance of the eligibility document that supports the official health certificate, the date of departure of the consignment and the date of signing of the official health certificate. New Zealand shall inform the border inspection post of arrival of any certification problem after departure from New Zealand.	

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	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Re-exports of imported animal products	96/93/EC	Animal Products Act 1999 Food Act 1981 Biosecurity Act 1993	Yes (1)	Animal products may be derived or partly derived from complying animal product(s) which originated in a third country/countries and establishment(s) eligible for trade with the EU and New Zealand.		Animal Products Act 1999 Food Act 1981 Biosecurity Act 1993	96/93/EC	Yes (1)	Animal products may be derived or partly derived from complying animal product(s) which originated in a third country/countries and establishment(s) eligible for trade with the EU and New Zealand.	
Microbiological monitoring/test system (1) (2) including: test methods, standards for sampling and preparation, and regulatory actions	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 2073/2005	Animal Products Act 1999	Yes (1)			Animal Products Act 1999	Regulations (EC) No 852/2004 (EC) No 853/2004 (EC) No 854/2004 (EC) No 2073/2005	Yes (1)		

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	EU Exports to New Zealand[1]					New Zealand Exports to EU				
	Trade conditions		Equivalence	Special conditions	Action	Trade conditions		Equivalence	Special conditions	Action
	EU standards	NZ standards				NZ standards	EU standards			
Establishment Listing Systems ([3])	Regulations (EC) No 178/2002 (EC) No 882/2004 (EC) No 852/2004 (EC) No 854/2004	Animal Products Act 1999	Yes (1)			Animal Products Act 1999	Regulations (EC) No 178/2002 (EC) No 882/2004 (EC) No 852/2004 (EC) No 854/2004	Yes (1)	Equivalence status applies to all animal product commodities accorded public health equivalence 'Yes (1)' as laid down in this Annex.	Procedures for establishment listings for non 'Yes (1)' commodities to be reviewed.

28. **Miscellaneous certification provisions: Attestations are to appear on the public or animal health certificate.**

Issue	Certification provisions
Q-fever	New Zealand is recognised as free of Q-fever. **For trade from the EU to NZ** in bovine semen and embryos, the Member State competent authority shall certify that: To the best of my knowledge and as far as I can ascertain, the donors have never been confirmed positive for Q-fever; AND For bovine semen EITHER The donors were subjected to a complement fixation test (CFT) (negative being no fixation of complement at dilution of 1:10 or higher) or ELISA test for Q-fever, on a sample collected between 21 to 120 days after each semen collection period (a period of 60 days or less) for export to New Zealand, with negative results.

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Issue	Certification provisions
	OR An aliquot of semen from each collection for export to New Zealand was tested using a laboratory validated Q-fever PCR test which is in accordance with the methods described in the Q-fever Chapter of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals. AND For bovine embryos EITHER The donors were subjected to a complement fixation test (CFT) (negative being no fixation of complement at dilution of 1:10 or higher) or ELISA test for Q-fever, on a sample collected between 21 to 120 days after each embryo collection period for export to New Zealand, with negative results. OR A sample of embryos/oocytes and collection and/or washing fluids from each collection for export to New Zealand was/were tested using a laboratory validated Q-fever PCR test which is in accordance with the methods described in the Q-fever Chapter of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.
BVD type II	New Zealand is recognised as free of Bovine viral diarrhoea virus (BVDV): Type II. **For trade from the EU to NZ** in bovine embryos, the Member State competent authority shall certify that: EITHER The donor animal was subjected to an antigen detection ELISA or virus isolation test for BVDV, with a negative result within thirty (30) days prior to entry into the herd of origin and has been in the herd of origin for more than six (6) months prior to embryo collection for this consignment and has remained isolated from other animals that have not been tested negative. OR From the first embryo collection taken from the donor animal for this consignment, either a pooled sample of non-viable oocytes/embryos and washing fluid (as per the OIE Code Chapter for *in vivo* derived embryos) or an embryo, has been subject to either virus isolation test or PCR test for BVDV with negative results.

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Issue	Certification provisions
Bluetongue	New Zealand is recognised as free of Bluetongue and Epizootic Haemorrhagic Disease. **For trade from the EU to NZ** in bovine semen, the Member State competent authority shall certify that: The bovine semen complies with the provisions of the Bluetongue Chapter of the OIE Code *mutatis mutandis*.
IBR	**For trade in live bovine animals from NZ** to Member States or regions thereof listed in Annex I to Decision 2004/558/EC New Zealand shall certify in accordance with Article 2 of Commission Decision 2004/558/EC, and to Member States or regions thereof listed in Annex II to Decision 2004/558/EC New Zealand shall certify in accordance with Article 3 of Decision 2004/558/EC. This attestation shall appear on the health certificate according to Commission Regulation (EU) No 206/2010.
BSE	**EU exports of products containing bovine, ovine or caprine materials to NZ** (in addition to full compliance with all other relevant EU standards) This product does not contain and is not derived from bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in the European Union and which has been produced in full compliance with Regulations of the European Parliament and the Council (EC) No 999/2001 and (EC) No 1069/2009 as applicable. Note: Products which contain bovine, ovine or caprine materials other than from those derived from animals born, continuously reared and slaughtered in the European Union are required to have that component certified in accordance with the relevant, additional third country provisions in the applicable NZ certification decision.
BSE	**NZ exports of products containing bovine, ovine or caprine materials to the EU** For human consumption — fresh meat, minced meat and meat preparations, meat products, treated intestines, rendered animal fats, greaves, and gelatine: (a) The country or region is classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a negligible BSE risk; (b) The animals from which the products of bovine, ovine and caprine animal origin were derived were born, continuously reared and slaughtered in a country with a negligible BSE risk. For by-products — rendered fats, pet food, blood products, processed animal protein, bones and bone products, category 3 material, and gelatine: The animal by-product does not contain and is not derived from bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001.

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Issue	Certification provisions
PRRS	**For trade from the EU to NZ in pig meat**, the Member State competent authority shall certify that: EITHER (i) derived from animals that were continuously resident since birth in Finland or Sweden, which is free of Porcine Reproductive and Respiratory Syndrome; OR (ii) cooked to one of the following core temperature/times: 56 degrees Celsius for 60 minutes; 57 degrees Celsius for 55 minutes; 58 degrees Celsius for 50 minutes; 59 degrees Celsius for 45 minutes; 60 degrees Celsius for 40 minutes; 61 degrees Celsius for 35 minutes; 62 degrees Celsius for 30 minutes; 63 degrees Celsius for 25 minutes; 64 degrees Celsius for 22 minutes; 65 degrees Celsius for 20 minutes; 66 degrees Celsius for 17 minutes; 67 degrees Celsius for 15 minutes; 68 degrees Celsius for 13 minutes; 69 degrees Celsius for 12 minutes; or 70 degrees Celsius for 11 minutes;

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Issue	Certification provisions
	OR (iii) cured where the product has been subjected to a procedure which ensures the meat meets one of the following requirements: reached a pH of 5 or lower; or was fermented (lactic curing) to a pH of 6,0 or lower and age-cured/ripened for at least 21 days; or qualified for official certification as Prosciutto di Parma or an equivalent 12 month curing process; OR (iv) prepared as consumer-ready cuts packaged for direct retail sale, not including minced (ground) meat, not including the head and neck, not exceeding 3 kg per package, with the following tissues removed: axillary, medial and lateral iliac, sacral, iliofemoral (deep inguinal), mammary (superficial inguinal), superficial and deep popliteal, dorsal superficial cervical, ventral superficial cervical, middle superficial cervical, gluteal and ischiatic lymph nodes; and any other macroscopically visible lymphatic tissue (i.e. lymph nodes and lymphatic vessels) encountered during processing; OR (v) none of the above (Note: These products need to be processed in New Zealand prior to being given a biosecurity clearance.)
Aujeszky's disease	**For trade in live pigs from NZ** to Member States or regions thereof listed in Annex I and Annex II to Decision 2008/185/EC New Zealand shall certify in accordance with Decision 2008/185/EC. This attestation shall appear on the health certificate according to Commission Regulation (EU) No 206/2010.
CSF **— feral pigs only**	**For trade from the EU to NZ** the Member State competent authority shall certify that the products were derived from areas free from CSF in the feral porcine population for the preceding 60 days. This attestation shall appear on the health certificate: ‘The product herein described was derived from wild pigs which were sourced from areas free from classical swine fever in the feral porcine population for the preceding 60 days.’

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Issue	Certification provisions
Live bees/ bumble bees	**For trade from NZ to the EU**, the health certificate(s) for live bees/bumble bees shall bear the following attestation: The bees/bumble bees ([1]), herein described: (a) come from a breeding apiary, which is supervised and controlled by the competent authority; (b) in the case of honey bees, hives come from an area which is not subject to any restrictions associated with an occurrence of American foul brood (the period of prohibition has been continued for at least 30 days following the last recorded case and the date of which all hives within a radius of three kilometres have been checked by the competent authority and all infected hives burned or treated and inspected to the satisfaction of the said competent authority); (c) are from hives or come from hives or colonies (in the case of bumble bees), which were inspected immediately prior to dispatch (normally within 24 hours) and show no clinical signs or suspicion of disease including infestations affecting bees. The packaging material, queen cages, accompanying products and food are new and have not been in contact with diseased bees or brood-combs, and all precautions have been taken to prevent contamination with agents causing diseases or infestations of bees. ([1]) *Delete as appropriate*
Colours for sanitary stamps	Regulation (EC) No 1333/2008 prescribes the colours that could be used for sanitary stamps.
Salmonella	**For trade from NZ to Sweden and Finland** The health certificate(s) for live animals and animal products listed below, shall bear the appropriate attestation set out in the corresponding legislation, if they are imported for consignment to either Sweden or Finland: For table eggs for human consumption New Zealand shall certify in accordance with Commission Regulation (EC) No 1688/2005 For live poultry for slaughter New Zealand shall certify in accordance with Annex A to Council Decision 95/410/EC For breeding poultry New Zealand shall certify in accordance with Annex II to Commission Decision 2003/644/EC For day old chicks New Zealand shall certify in accordance with Annex III to Commission Decision 2003/644/EC

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Issue	Certification provisions
	For laying hens New Zealand shall certify in accordance with Annex II to Commission Decision 2004/235/EC For fresh meat covered by Regulation (EC) No 1688/2005, the following attestation is to be added 'The fresh meat has been subject to microbiological testing for salmonella as provided for in Regulation (EC) No 1688/2005 by sampling in the establishment of origin of this meat.'
Salmonids	**For trade from the EU to NZ** The consignment contains only beheaded, gilled, gutted and sexually immature Salmonids of the genera *Onchorhynchus, Salmo* or *Salvelinus*.
Eggs/roes	**For trade from the EU to NZ** Must be treated to render eggs/roe non-viable, commercially packaged and shelf stable.
Thermised cheeses	**For trade from the EU to NZ** The thermised cheese has a moisture content of less than 39 % and pH less than 5,6. The milk used to produce this cheese was rapidly heated to at least 64,5 °Celsius for 16 seconds. The cheese was stored at not less than 7° Celsius for 90 days.

29. **Mutually agreed disease control measures**

29.A. **Mutually agreed disease status for specific diseases**

Rabies	New Zealand, UK, Malta, Ireland and Sweden are recognised as free of rabies
Equine infectious anemia	New Zealand is recognised as free of EIA
Brucellosis	New Zealand is recognised as free of *Brucella abortus* and *B. mellitensis*
Q-fever	New Zealand is recognised as free of Q-fever
BVD type II	New Zealand is recognised as free BVD type II

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Issue	Certification provisions
Bluetongue and EHD	New Zealand is recognised as free of Bluetongue and EHD EU makes a submission to NZ for EHD freedom
Small hive beetle	New Zealand and the EU are recognised as free of small hive beetle
Tropilaelaps mite	New Zealand and the EU are recognised as free of tropilaelaps mite

No 29.B. **Mutually agreed disease control measures in the event of the occurrence of a specific disease**

Official health certificates shall, in accordance with Section 1(b) of Annex VII to the Agreement, bear the relevant additional attestation(s) listed under Chapter 29 of this Annex.

General Attestation for all commodities:

The commodity herein described was kept separate from all other commodities that did not meet the requirements during all stages of production, storage and transport and all necessary precautions were taken to prevent contamination of the commodity with any potential source of [insert relevant disease noted in the disease column below] virus.

Disease specific attestation:

The commodities listed in Chapter 29 i) to xxx) shall in addition to the General Attestation (noted above) for all commodities bear the relevant disease attestation(s) below:

Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(i) **Milk and Milk Products:** **8.0 13.0**	FMD	The milk/milk products herein described: EITHER 1*) have undergone sterilisation of at least $F_0 3$. OR 2*) have undergone an ultra-high temperature (UHT) treatment at 132 °C for at least 1 second.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
		OR 3*) had a pH of less than 7,0 prior to undergoing a high temperature — short time pasteurisation (HTST) treatment at 72 °C for 15 seconds. OR 4*) had a pH of more than 7,0 prior to undergoing a double high temperature — short time pasteurisation (HTST) treatment at 72 °C for 15 seconds. OR 5*) have been subjected to high temperature — short time pasteurisation (HTST) combined with the lowering of pH below 6 for one hour. OR 6*) have been subjected to high temperature — short time pasteurisation (HTST) combined with additional heating to 72 °C combined with desiccation or an equivalent validated and approved drying/desiccation process that achieves at minimum an equivalent thermal effect of 72 °C for 15 seconds.
(ii) **Meat (including minced meat) and meat preparations from bi-ungulates excluding head, feet, viscera and meat from swine (*suidae*):** **4.A 4.C 5.A 5.C**	FMD	The [*insert relevant commodity*] herein described (excluding feet, head and viscera) was: 1. derived from animals that have been subjected to *ante-mortem* and *post-mortem* inspections and have been found free of any sign suggestive of FMD; 2. derived from deboned carcasses from which the offal and major lymphatic glands have been removed; 3. subject to maturation at a temperature above + 2 °C for at least 24 hrs and reached a pH value of below 6 when tested in the middle of the *longissimus dorsi* muscle after maturation and before deboning; 4. was not derived from animals slaughtered or processed in an establishment located within a designated protection or surveillance zone, 5. Meat sourced from animals within the protection and surveillance zones is subject to official control and has been identified and controlled so as to ensure its exclusion from this consignment.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(iii) **Meat (including minced meat) and other animal products (including offal) derived from bi-ungulates including swine (*suidae*):** **4.A 4.C 5.A 5.C 7.A 7.B 7.C 7.D 7.E 7.F 7.G 11.E**	FMD	The [*insert relevant commodity*] herein described was: I) derived from animals that have been subjected to *ante-mortem* and *post-mortem* inspection and have been found free of any sign suggestive of FMD; AND EITHER 2*) derived from animals slaughtered 21 days prior to the estimated date of earliest infection in the territory; and not derived from animals slaughtered or processed in an establishment located within a designated protection or surveillance zone. OR 3*) derived from animals that have been resident on a holding for at least 21 days and were identified so as to allow the tracing of the holding of origin; but not derived from animals resident in holdings within a protection or surveillance zone; and the commodity has been clearly identified and detained under official supervision for at least 7 days and was not released until any suspicion of infection with the foot-and-mouth disease virus on the holding of origin has been officially ruled out at the end of the detention period; AND 4. Meat sourced from animals within the protection and surveillance zones is subject to official control and has been identified and controlled so as to ensure its exclusion from this consignment.
(iv) **Meat and meat preparations from Poultry (including turkeys):** **4.B 4.C 5.B 5.C**	HPNAI — Notifiable in accordance with OIE Terrestrial Animal Health Code criteria	The [*insert relevant commodity*] herein described was derived from animals that: EITHER 1*) have been sourced from a holding situated outside a protection or a surveillance zone; and all meat sourced from animals within the protection and surveillance zones is subject to official control and has been identified and controlled so as to ensure its exclusion from this consignment.; OR 2*) have been sourced from a holding within a surveillance zone but outside a protection zone and have been tested to give a 95 % probability of detecting a 5 % prevalence of HPNAI infection not more than 7 days prior to slaughter using virus detection tests, and/or serological tests, with negative results; and have been slaughtered in a designated establishment which has not processed poultry infected with HPNAI since last cleaned and disinfected, and have been subjected to *ante-mort*em and *post-mortem* examinations and have shown no signs suggestive of HPNAI; and all meat sourced from animals within the protection zone is subject to official control and has been identified and controlled so as to ensure its exclusion from this consignment.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
		OR 3*) were processed on a date at least 21 days before the estimated date of earliest infection.
(v) **Meat and meat preparations from Poultry (including turkeys):** **4.B 4.C 5.B 5.C**	LPNAI Notifiable in accordance with OIE Terrestrial Animal Health Code criteria	The [*insert relevant commodity*] herein described was derived from animals that; 1. have been sourced from a holding in which there has been no evidence of LPNAI during the past 21 days; 2. have been slaughtered in an approved establishment which has not processed poultry infected with LPNAI since last cleaned and disinfected; 3. have been subjected to *ante-mortem* and *post-mortem* examinations and have shown no signs suggestive of LPNAI;
(vi) **Meat and meat preparations from Poultry (including turkeys):** **4.B 4.C 5.B 5.C**	ND	The [*insert relevant commodity*] herein described was derived from: 1. Animals from holdings free from ND and not situated in an ND protection or surveillance zone; AND EITHER 2*) have not been vaccinated against ND; OR 3*) were vaccinated against ND using a vaccine complying with the standards described in Commission Decision 93/152/EEC (the nature of the vaccine used and the date of vaccination shall also be stated in the certificate). AND 4. The animals showed no clinical sign of ND on the day of shipment to the slaughter house and were further subjected to *ante-mortem* and *post-mortem* examination and having showed no clinical signs suggestive of ND; were slaughtered in an approved establishment that is subject to regular inspection by the Veterinary Competent Authority and which has not processed poultry infected with ND since having last cleaned and disinfected.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(vii) **Meat Products and other processed products derived from bi-ungulates including swine (*suidae*) and poultry (including turkeys):** **6.A 6.B 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	FMD, CSF, SVD, ASF, RND, ND, LPNAI, HPNAI, PPR	The [*insert relevant commodity*] herein described has been heat treated in a hermetically sealed container with an F_0 value of 3,00 or more
(viii) **Meat Products and other processed products derived from bi-ungulates including swine (*suidae*) and poultry (including turkeys):** **6.A 6.B 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	FMD, CSF, SVD, RND, ND, LPNAI, HPNAI, PPR	EITHER 1*) The [*insert relevant commodity*] herein described has been heat treated to a minimum temperature of 70 °C throughout the product. OR 2*) The [*insert relevant commodity*] herein described has been heat treated to 70 °C for minimum 30 minutes or an equivalent validated and approved thermal process.

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Commodity		Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(ix)	**Meat Products and other processed products derived from bi-ungulates including swine (*suidae*):** **6.A 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	FMD, CSF, SVD, ASF, RND, PPR	The [*insert relevant commodity*] herein described has undergone heat treatment in a hermetically sealed container to at least 60 °C for a minimum of 4 hours, during which time the core temperature has reached at least 70 °C for 30 minutes.
(x)	**Meat Products and other processed products derived from swine (*suidae*):** **6.A 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	ASF	The [*insert relevant commodity*] herein described has been heat treated to a minimum temperature of 80 °C throughout the product.

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Commodity		Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(xi)	**Meat Products and other processed products (boneless) derived from bi-ungulates including swine (*suidae*):** **6.A 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	FMD, CSF, SVD, ASF, RND	The [*insert relevant commodity*] herein described is boneless and has undergone a natural fermentation and maturation process for not less than nine months resulting in the following characteristics: Aw value of not more than 0,93 or a pH value of not more than 6,0.
(xii)	**Meat Products and other processed products (including bone in) derived from bi-ungulates including swine (*suidae*):** **6.A 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	FMD, CSF, SVD	The [*insert relevant commodity*] herein described which may contain bone has undergone a natural fermentation and maturation process for not less than nine months resulting in the following characteristics: Aw value of not more than 0,93 or a pH value of not more than 6,0.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(xiii) **Meat Products and other processed products derived from bi-ungulates including swine (*suidae*):** **6.A 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	FMD, PPR	The [*insert relevant commodity*] herein described has been heat treated to ensure a core temperature of at least 65 °C is reached for the time necessary to achieve a pasteurisation value (PV) equal to, or more than, 40.
(xiv) **Meat Products and other processed products derived from swine (*suidae*):** **6.A 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	CSF	The [*insert relevant commodity*] herein described is a dry cured pork meat[1] and is; EITHER 1*) Bone-in Italian style pork ham that has been cured with salt and dried for a minimum of 313 days[1]; OR 2*) Bone-in Spanish style pork, Iberian shoulder, that has been cured with salt and dried for a minimum of 252 days[1]; OR 3*) Bone-in Spanish style pork, Iberian loin, that has been cured with salt and dried for a minimum of 126 days[1]; OR 4*) Bone-in Spanish style pork, Serrano ham, that has been cured with salt and dried for a minimum of 140 days[1]. Foot note [1]: At the time of publication import conditions for pork meat into New Zealand may apply curing times exceeding the minimum stated for CSF.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(xv) **Meat Products and other processed products derived from swine (*suidae*):** **6.A 6.C 6.D 7.B 7.C 7.D 7.E 7.F 7.G7.H**	ASF	The [*insert relevant commodity*] herein described has been subject to treatment involving natural fermentation and maturation during at least 190 days for hams and 140 days for loins.
(xvi) **Animal Casings derived from ruminants:** **7.A 12.0**	FMD	The animal casings herein described have been cleaned, scraped and either salted with sodium chloride for 30 days or bleached or dried after scraping and were protected from recontamination after treatment.
(xvii) **Processed (rendered) Animal Protein, lards, fats and petfood derived from ungulates and poultry (including turkeys):** **15.0 17.0 21.0**	FMD, SVD, RND, PPR, ASF, ND, LSD	The [*insert relevant commodity*] herein described has been thermally treated in accordance with minimum regulatory standards and to a minimum 90 degrees Celsius for ten minutes throughout the product.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(xviii) **Wool and fibre from ruminants:** **20.0**	FMD, RND	EITHER 1*) The [*insert relevant commodity*] herein described has been stored at 18 °C for 4 weeks, or 4 °C for 4 months, or 37 °C for 8 days OR 2*) The [*insert relevant commodity*] herein described has been subject to industrial scouring by immersion in water soluble detergent at 60-70 °C. OR 3*) The unprocessed [*insert relevant commodity*] has been cleaned, dried and securely enclosed in packaging in accordance with the requirements of Regulation (EC) No 1069/2009
(xix) **Treated Hides and Skins:** **19**	FMD, RND	The hides or skins herein described have been salted for 7 days in sea salt containing at least 2 % sodium carbonate.
(xx) **Treated Hides and Skins:** **19**	FMD	EITHER 1*) The hides or skins herein described have been dry or wet salted for 14 days prior to dispatch and have been shipped by sea. OR 2*) The hides and skins herein described have been dried for 42 days at temperatures of at least 20 °C.
(xxii) **Fully Treated Hides and Skins (wet blue, pickled, limed or hides that have completed the tanning process):** **19**	FMD, RND	Fully treated hides and skins can be traded without restriction provided that these products have been submitted to the usual chemical and mechanical processes in use in the tanning industry. The following attestation may be applied in order to facilitate trade: The fully treated hides and skins described have been submitted to the usual chemical and mechanical processes in use in the tanning industry.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(xxiii) **Bovine Semen:** **1**	FMD	The semen herein described: EITHER 1*) was derived from donor animals which were kept in a semen collection centre where no animals have been added in last the 30 days before collection and FMD has not occurred within 10 kilometres for 30 days before and after collection, and which showed no clinical sign of FMD on the day of collection, have not been vaccinated against FMD and were subjected, not less than 21 days after collection of the semen, to a tests for antibodies against FMD virus, with negative results and no other animals present in the semen collection centre has been vaccinated against FMD. Additionally, the semen was collected from a semen collection centre not located within a protection or surveillance zone and any semen collected within a protection and surveillance zone has been clearly identified and detained under official supervision; and the semen collected, was further processed and stored in conformity with the provisions of Chapter 4.5 or Chapter 4.6 of the OIE Terrestrial Animal Health Code as relevant and was further stored in the country of origin for a period of at least one month following collection, and during this period no animal on the establishment where the donor animals were kept showed any sign of FMD. OR 2*) has been collected and stored frozen at least 21 days before the estimated date of earliest infection with the foot-and-mouth disease virus on a holding in the protection and surveillance zone; and any semen collected after the date of earliest infection has been stored separately and was only released after all the measures relating to the outbreak of FMD have been removed; and all animals accommodated in the semen collection centre have undergone a clinical examination and samples taken have been subjected to a serological test to substantiate the absence of infection in the centre concerned; and the donor animals have been subjected with negative result to a serological test for the detection of antibodies against the FMD virus on a sample taken not earlier than 28 days after the collection of the semen.
(xxiv) **Bovine Semen:** **1**	BT	The semen herein described was derived from donor animals: EITHER That were kept in a vector-protected establishment for at least 60 days before the commencement of, and during, collection of the semen; OR That were subjected to a serological test according to the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals to detect antibody to the BTV group, with negative results, at least every 60 days throughout the collection period and between 21 days and 60 days after the final collection for the consignment to be exported:

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
		OR That were subjected to an agent identification test according to the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals on blood samples collected at commencement and conclusion of, and at least every 7 days (virus isolation test) or at least every 28 days (for PCR test) during, semen collection for the consignment to be exported, with negative results: OR The semen collection centre is not within an infected (restricted) zone. Semen from infected (restricted) zones has been clearly identified and detained under official supervision. AND The semen was collected, processed and stored in conformity with the OIE standards.
(xxv) **Bovine Semen:** **1**	LSD	The semen herein described was derived from donor animals: That showed no clinical sign of LSD on the day of collection of the semen and for the following 28 days; and the animals were kept in the exporting country for the 28 days prior to collection, in a semen collection centre where no case of LSD was officially reported during that period, and the centre was not situated in either a LSD infected zone or buffer zone and any semen from buffer zone has been clearly identified and controlled.
(xxvi) ***In vivo* derived bovine embryos (except embryos that have been subjected to penetration of the *zona pellucida*):** **2**	FMD	The *in vivo* derived embryos herein described were derived from donors that: Were free of clinical signs of FMD, at the time of collection; and from which the embryos were conceived by artificial insemination using semen collected, processed and stored in semen collection centres approved by the competent authority in conformity with OIE standards. In addition the embryos have been collected, processed and stored in accordance with standards laid down by the competent authority; AND The donor animals from which the embryos were collected originate from a herd(s) that was/were not located within a protection or surveillance zone. Embryos collected within the protection and surveillance zones have been clearly identified and detained under official supervision.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(xxvii) ***In vivo* derived bovine embryos (except embryos that have been subjected to penetration of the *zona pellucida*):** **2**	BT	The *in vivo* derived embryos herein described were derived from donors that: Were free of clinical signs of BT at the time of collection and from which the embryos were conceived by artificial insemination using semen collected, processed and stored in semen collection centres approved by the competent authority in conformity with the OIE standards. AND The embryos were collected, processed and stored in accordance with standards laid down by the competent authority.
(xxviii) ***In vivo* derived bovine embryos (except embryos that have been subjected to penetration of the *zona pellucida*):** **2**	VS	The *in vivo* derived embryos herein described were derived from donors that: Were kept for 21 days prior to, and during, collection in an establishment where no case of VS was reported during that period and were subject to a diagnostic test for VS, with negative results, within 21 days prior to embryo collection. In addition the embryos were collected, processed and stored in conformity with OIE notified standards; and the establishment was not located within a protection or surveillance zone. Embryos collected within protection and surveillance zones has been clearly identified and detained under official supervision.
(xxix) ***In vivo* derived bovine embryos (except embryos that have been subjected to penetration of the *zona pellucida*):** **2**	CBPP	The *in vivo* derived embryos herein described were derived from donors that: EITHER 1*) have not been vaccinated against CBPP and were subjected to the complement fixation test for CBPP with negative results, on two occasions, with an interval of not less than 21 days and not more than 30 days between each test, the second test being performed within 14 days prior to collection; and were isolated from other domestic *bovidae* from the day of the first complement fixation test until collection; OR 2*) were vaccinated using a vaccine complying with the standards described in the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals not more than 4 months prior to collection; AND showed no clinical sign of CBPP on the day of collection of the embryos; and were kept since birth, or for the past 6 months, in a herd(s) where no case of CBPP was reported during that period, and that the herd(s) was/were not situated in a CBPP infected zone; and the embryos were collected, processed and stored in accordance with standards laid down by the competent authority.

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Commodity	Disease	Disease Attestation Number*) Optional attestations need only be applied to the certificate when applicable.
(xxx) **Poultry hatching eggs:** **2**	LPNAI, HPNAI — Notifiable in accordance with OIE Terrestrial Animal Health Code criteria Avian influenza (OIE notifiable) Newcastle disease	**For trade from the EU to NZ:** The poultry hatching eggs herein described were derived from source flocks and hatcheries within a Ministry for Primary Industries approved compartment free of notifiable avian influenza [and/or] Newcastle disease [*delete as appropriate*]
(xxxi) **Live bees/ bumble bees:** **3**	Small hive beetle (*Aethina tumida*)	**For trade from NZ to the EU:** (a) hives come from an area at least 100 km radius which is not subject to any restrictions associated with the suspicion or confirmed occurrence of the small hive beetle (*Aethina tumida*) and where these infestations are absent; (b) the bees/bumble bees ([1]) as well as their packaging have undergone a visual examination to detect the occurrence of the small hive beetle (Aethina tumida) or their eggs and larvae. ([1]) *Delete as appropriate*
(xxxii) **Live bees/ bumble bees:** **3**	Tropilaelaps mite (*Tropilaelaps* spp.)	**For trade from New Zealand to the EU:** (a) hives come from an area at least 100 km radius which is not subject to any restrictions associated with the suspicion or confirmed occurrence of the Tropilaelaps mite (*Tropilaelaps* spp.), and where these infestations are absent; (b) the bees/bumble bees ([1]) as well as their packaging have undergone a visual examination to detect the occurrence of the Tropilaelaps mite (*Tropilaelaps* spp.). ([1]) *Delete as appropriate*

([1]) For exported products it is the responsibility of the exporter (food business operator) to ensure that exported products meet the microbiological food safety criteria of the importing party.
([2]) Applies to the meat, fishery and dairy sectors.
([3]) New Zealand establishments and facilities information will be entered into the EU TRACES system (or any successor system) by the New Zealand Competent Authority. New Zealand guarantees that the establishments fulfil the conditions as laid down in the Agreement. The Commission will update and publish the information on the Commission website without undue delay and normally within 2 working days. The Commission may, where a guarantee is unsatisfactory, not publish an establishment on the Commission website. If the Commission decides not to publish an establishment on the Commission website it will provide the reason(s) to the New Zealand authority without undue delay.

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ANNEX VI

GUIDELINES ON PROCEDURES FOR CONDUCTING AN AUDIT

For the purposes of this appendix 'audit' means assessment of performance.

1. **General principles**

1.1. Audits should be made in cooperation between the auditing party (the 'auditor') and the audited party, (the 'auditee') in accordance with the provisions set out in this Annex. Checks of establishments or facilities may be made as considered necessary.

1.2. Audits should be designed to check the effectiveness of the controlling authority rather than to reject individual animals, groups of animals, consignments of food or establishments. Where an audit reveals a serious risk to animal or human health, the auditee shall take immediate corrective action. The process can include study of the relevant regulations, method of implementation, assessment of the end result, level of compliance and subsequent corrective actions.

1.3. The frequency of audits should be based on performance. A low level of performance should result in an increased frequency of audit; unsatisfactory performance must be corrected by the auditee to the auditor's satisfaction.

1.4. Audits, and the decisions based on them, shall be made in a transparent and consistent manner.

2. **Principles relating to the auditor**

Those responsible for conducting the audit should prepare a plan, preferably in accordance with recognized international standards, that covers the following points:

2.1. the subject, depth and scope of the audit;

2.2. the date and place of the audit, along with a timetable up to and including the issue of the final report;

2.3. the language or languages in which the audit will be conducted and the report written;

2.4. the identity of the auditors including, if a team approach is used, the leader. Specialized professional skills may be required to carry out audits of specialized systems and programmes;

2.5. a schedule of meetings with officials and visits to establishments or facilities, as appropriate. The identity of establishments or facilities to be visited need not be stated in advance;

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2.6. subject to provisions on freedom of information, respect of commercial confidentiality shall be observed by the auditor. Conflicts of interest must be avoided;

2.7. respect of the rules governing occupational health and safety, and the rights of the operator.

This plan should be reviewed in advance with representatives of the auditee.

3. **Principles relating to the auditee**

The following principles apply to actions taken by the auditee, in order to facilitate audit.

3.1. The auditee must cooperate fully with the auditor and should nominate personnel responsible for this task. Cooperation may include, for example:

— access to all relevant regulations and standards,

— access to compliance programmes and appropriate records and documents,

— access to audit and inspection reports,

— documentation concerning corrective actions and sanctions,

— facilitating entry to establishments.

3.2. The auditee must operate a documented programme to demonstrate to third parties that standards are being met on a consistent and uniform basis.

4. **Procedures**

4.1. *Opening meeting*

An opening meeting should be held between representatives of both parties. At this meeting the auditor will be responsible for reviewing the audit plan and confirming that adequate resources, documentation, and any other necessary facilities are available for conducting the audit.

4.2. *Document review*

The document review may consist of a review of the documents and records referred to in paragraph 3.1, the structures and powers of the auditee, and any relevant changes to food inspection and certification systems since the adoption of this Agreement or since the previous audit, with emphasis on the implementation of elements of the system of inspection and certification for animals or products of interest. This may include an examination of relevant inspection and certification records and documents.

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4.3. *On-site verification*

4.3.1. The decision to include this step should be based on a risk assessment, taking into account factors such as the animals or products concerned, the history of conformity with requirements by the industry sector or exporting country, the volume of product produced and imported or exported, changes in infrastructure and the nature of the national inspection and certification systems.

4.3.2. On-site verification may involve visits to production and manufacturing facilities, food-handling or storage areas and control laboratories to check on compliance with the information contained in the documentary material referred to in 4.2.

4.4. *Follow-up audit*

Where a follow-up audit is being conducted in order to verify the correction of deficiencies, it may be sufficient to examine only those points which have been found to require correction.

5. **Working documents**

Forms for reporting audit findings and conclusions should be standardized as much as possible in order to make the approach to audit more uniform, transparent and efficient. The working documents may include any checklists of elements to evaluate. Such checklists may cover:

— legislation,

— structure and operations of inspection and certification services,

— establishment details and working procedures,

— health statistics, sampling plans and results,

— compliance action and procedures,

— reporting and complaint procedures,

— training programmes.

6. **Closing meeting**

A closing meeting must be held between representatives of both parties, including, where appropriate, officials responsible for the national inspection and certification programmes. At this meeting the auditor will present the findings of the audit. The information should be presented in a clear, concise manner so that the conclusions of the audit are clearly understood.

An action plan for correction of any deficiencies noted should be drawn up by the auditee, preferably with target dates for completion.

7. **Report**

The draft report of the audit shall be forwarded to the auditee as soon as possible. The auditee shall have one month in which to comment on the draft report; any comments made by the auditee shall be included in the final report.

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ANNEX VII

CERTIFICATION

Official health certificates will cover consignments of live animals and/or animal products being traded between the Parties.

Section 1: Health attestations:

(a) For commodities with equivalence 'Yes-1' agreed

(i) The following model health attestation to be used (equivalence for animal and/or public health as appropriate). Refer Yes (1) Annex V;

'The live animal(s) or animal product(s) herein described, complies/y with the relevant (European Union/New Zealand (*)) standards and requirements which have been recognized as equivalent to the (New Zealand/European Union (*)) standards and requirements as prescribed in the European Union/New Zealand Agreement on sanitary measures (Council Decision 97/132/EC).

Specifically, in accordance with (insert … exporting Party's legislation) (**)

(*) Delete as appropriate.
(**) Optional, at the discretion of the importing Party.'

AND

(ii) The additional attestation(s) described in Chapter 28 of Section 5 of Annex V, as relevant and referred to as 'Special Conditions' within Annex V, to be used.

(iii) For EU exports to New Zealand, the additional attestation(s) to be used: 'the animal product is eligible for intra-Union trade without restriction'.

(iv) For exports from New Zealand: For consignments of commodities for which the model health attestation as referred to in Section 1 paragraph (a)(i) is prescribed and equivalence is established in Annex V, Section 5, Chapter 28, Subchapter 'Certification systems', the additional attestation to be used when certificates are issued after the date of departure ([13]) of consignments: 'The undersigned officer certifies this consignment on the basis of eligibility document(s) (specify reference to the appropriate Eligibility Document(s) ED)) issued on (insert date), which were ascertained by him/her and were issued prior to the departure of the consignment'.

(b) For all commodities

Following confirmation by the exporting party, in accordance with Article 12, that a disease listed in Chapter 29.B. of Section 5 of Annex V has occurred, the relevant additional attestation(s), as described in Chapter 29.B. of Section 5 of Annex V, shall be applied to official health certificates. The relevant additional attestation(s) provided for in Chapter 29.B. of Section 5 of Annex V to be used until a regionalisation decision is taken by the exporting party, in accordance with Article 6, or as otherwise mutually agreed.

Section 2: Completion of Certificates:

(a) When issuing a paper certificate, the signature and official seal applied must be in a colour different to that of the printing.

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(b) For exports from New Zealand: when a paper official health certificate is issued, the official health certificate shall be issued in English, as well as in one of the languages of the Member State in which the border inspection post where the consignment is presented, is situated.

(c) For exports from the European Union: the official health certificate shall be issued in the language of the Member State of origin, as well as in English.

(d) Each consignment intended for export shall be supported by an original health certificate(s), or original veterinary document(s) or other original document(s) where specified under the Agreement which convey agreed sanitary information.

(e) Minor modifications to the format of the model certificate are permitted.

(f) The official health certificates do not need to include the explanatory notes providing guidance for completion, nor the attestations that are irrelevant to the consignment.

Section 3: Electronic Data Transmission:

(a) The exchange of original veterinary certificate(s) or other original document(s)/information may occur by paper based systems and/or secure methods of electronic data transmission offering equivalent certification guarantees, including the use of digital signature and non-repudiation mechanism. Where the exporting Party elects to provide electronic official health certificates and/or veterinary document(s), the importing Party must have determined that equivalent security guarantees are being provided. The importing Party's agreement for the exclusive use of electronic certification can either be recorded in one of the Annexes to the Agreement or by correspondence in accordance with Article 16(1) to the Agreement. The Parties shall take all necessary steps to ensure the integrity of the certification process, to guard against fraud and prevent false and misleading certification.

Electronic data transmission systems offering equivalent guarantees:

New Zealand — E-cert

EU — TRACES

(b) The official health certificate shall be issued and provided to the border inspection post either:

(i) As an original signed paper certificate, or

(ii) Electronically through use of electronic data transmission using E-cert and TRACES according to the procedure described in Section 3 (a).

Section 4: Controls:

The controlling authority shall ensure that official certifying officers are aware of the importing party's health conditions as prescribed in this Agreement and are obliged to certify to these requirements where appropriate.

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ANNEX VIII

FRONTIER CHECKS AND INSPECTION FEES

A. FRONTIER CHECKS ON CONSIGNMENTS OF LIVE ANIMALS AND ANIMAL PRODUCTS

Type of frontier check [1]**:**	**Rate in %**
1. **Documentary and Identity checks**	100
Both Parties will perform documentary checks	
Identity check means a discretionary [2] confirmatory check by the Competent Authority to ensure that the sanitary certificate(s)/document(s) or other document(s) provided for by sanitary legislation correspond with the product within the consignment [3]. In the case of sealed containers, such identity check may consist of only verifying that the seals are intact and that container identity information and the seal number correspond to those given in the accompanying sanitary documentation or certificate.	
2. **Physical checks (including random or targeted)**	
Live animals, except bees and bumble bees	100
Queen bees and small colonies of bumble bees	100
Bees and bumble bees packages	50 [4]
Semen/embryos/ova	10
Live animals [5] and animal products for human consumption listed in Annex V to Council Decision 97/132/EC	1
Animal products not for human consumption listed in Annex V to Council Decision 97/132/EC	1
Processed animal protein not for human consumption (bulked)	100 % for the first 6 consignments and then 1-10 %.

[1] The Competent Authority may delegate these activities, including physical inspections, to a responsible person or an agency, in accordance with the legislation of the importing Party.

[2] In accordance with the legislation of the importing Party.

[3] For the purposes of this Annex, 'consignment' means a quantity of products of the same type, covered by the same veterinary certificate(s) or veterinary document(s), or other document(s) provided for by veterinary legislation, conveyed by the same means of transport and coming from the same third country or part of such country. 'Same means of transport' means carrier (e.g. vessel, aircraft).

[4] For consignments of packaged bees containing less than 130 packages 50 % of the consignment is to be subject to inspection. For consignments containing more than 130 packages, a sample of 65 packages randomly selected from the consignment must be inspected to achieve a 95 % confidence interval of detecting 5 % incidence of disease.

[5] As covered by Chapter 10 of Annex V.

▼M6

B. INSPECTION FEES

The fees specified in B.I and II of this Annex shall be applied to imports.

Fees, unless otherwise agreed, shall be set so that they only recover the actual costs of border inspection service and shall not be higher than the equivalent consignment fee charged for the same commodity imported from other third countries.

B.I. **For the European Union**

Live animals and germplasm inspection fees:

Inspection fees shall be applied in accordance with Annex V to Regulation (EC) No 882/2004.

Products of animal origin:

Inspection fees shall be applied in accordance with Annex V to Regulation (EC) No 882/2004 with a reduction of 22,5 % ([1]). However, for the transit of goods through the Union, inspection fees shall be applied in accordance with Annex V to Regulation (EC) No 882/2004 without reduction.

B.II. **For New Zealand**

Live animals and germplasm inspection fees:

Inspection fees shall be applied in accordance with New Zealand — Biosecurity (Costs) Regulations.

Products of animal origin:

Documentary and identity checks inspection fees:

Single consignment — Maximum 149,60(+ gst) NZD per consignment

Multi container consignments — Maximum 149,60(+ gst) NZD for the first container and a Maximum of 75 (+ gst) NZD/container for additional containers

Break bulk consignments — Maximum 149,60(+ gst) NZD/hour

Documentary, identity + physical checks inspection fees:

Single consignment — inspection fees applied in accordance with New Zealand regulations:

Animal Health Biosecurity (Costs) Regulations

Public Health Fees and Charges Regulation

([1]) Calculated on the assumption that the rate of the physical checks for New Zealand imports is only 10 % of the normal physical checks rate applied to other third countries and assuming that the costs for physical checks account for 25 % of the total fees costs.

▼M6

Inflation adjustment for New Zealand's inspection fees:

New Zealand inspection fees may be adjusted on an annual basis in accordance with the following formula:

Maximum inspection fee =

Annex VIII listed inspection fee × (1 + average inflation rate/100*)(Current year — 2009)

* as calculated on an ongoing basis for New Zealand as published by the Reserve Bank of New Zealand.

▼B

ANNEX IX

OUTSTANDING ISSUES

— Provision of electronic access to draft standards.

— Conditions for live animals and animal products transiting through the territories of the Parties to this Agreement.

— Consideration of the inclusion of other species in the manufacture of lards and fats (e.g. poultry).

— Trade conditions for packaged raw petfood intended for direct sale to the consumer.

— Trade conditions for cervine velvet.

— Progress towards implementing export health certificate transfer from controlling authority to controlling authority using the electronic data interchange system (EDI) (utilizing the established UN/Edifact and Sancrt protocols).

▼M3

ANNEX X

CONTACT POINTS

For New Zealand

Counsellor (Veterinary and Technical)
New Zealand Mission to the European Union
Square De Meeus/De Meeusplein 1
B-1000 Brussels
Tel. (0032) 25 50 12 19
Fax (0032) 25 13 48 56

For the European Community

The Director
Directorate E: Food Safety: plant health, animal health and welfare, international questions
Directorate-General Health and Consumer Protection
European Commission
Rue Froissart/Froissartstraat 101, Room 8/58
B-1040 Brussels
Tel. (0032) 22 95 68 38
Fax (0032) 22 96 42 86