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DEPARTMENT OF AGRICULTURE

DISEASES OF ANIMALS (NORTHERN IRELAND) ORDER 1981

Landing of Carcases and Animal Products Order (Northern Ireland) 1985 (As Amended)

GENERAL IMPORT LICENCE AMENDMENT NOTICE

The following import licences, dated 8th April 1993, shall be amended as indicated below:

ARI.19 on the import of bovine semen from Canada;

ARI.20 on the import of bovine semen from Norway;

ARI.21 on the import of bovine semen from the United States of America;

ARI.22 on the import of bovine semen from Poland;

ARI.23 on the import of bovine semen from Hungary;

ARI.24 on the import of bovine embryos from Canada, New Zealand, United States of America;

ARI.25 on the import of bovine semen from Switzerland;

– any reference to the 'Importation of Semen, Embryos and Ova Order (Northern Ireland) 1993' shall be replaced by the 'Landing of Carcases and Animal Products Order (Northern Ireland) 1985 (as amended)'.

Catherine Coll,
Officer of the Department of
Agriculture for Northern Ireland.

Dated the 6th September, 1995.

DISEASES OF ANIMALS (NORTHERN IRELAND) ORDER 1981

Landing of Carcases and Animal Products Order (Northern Ireland) 1985 (As Amended)

Licence No.: DANI/GEN/95/22

GENERAL IMPORT LICENCE AMENDMENT NOTICE

The Department of Agriculture for Northern Ireland, in accordance with the terms of the above legislation, hereby authorises, subject to the conditions attached to this licence, the landing in Northern Ireland of:

Product: Products derived from ruminating animals or swine in sealed impervious containers for use as in-vitro laboratory or diagnostic reagents.

From: Australia, Canada, Falkland Islands, Iceland, Japan, New Zealand, Norway, United States of America:

until further notice or unless the licence is revoked by the Department.

Catherine Coll,
Officer of the Department of
Agriculture for Northern Ireland.

Dated the 6th September, 1995.

CONDITIONS ATTACHED TO THIS LICENCE

1. The products must be shipped in sealed, impervious containers.
2. The containers or their outside packaging must be clearly labelled "For in-vitro diagnostic use only" or "For in-vitro laboratory use only".
3. The products may be used in Northern Ireland only for in-vitro laboratory or diagnostic work and any product literature inserts must state that the products or their residues must not be allowed to come into contact with ruminating animals or swine.
4. Packaging, wrapping and any residual matter must be disposed of in such manner so that any animal health risk is eliminated.
5. Consignments imported directly into Northern Ireland must be:
   (a) imported through the border inspection post at Belfast Port or Belfast International Airport;
   (b) notified in advance to the official veterinarian at the border inspection post – one working day's advance notice must be given for products entering Belfast Port and at least 6 hours notice of arrival must be given during the working day of the border inspection post at Belfast International Airport;
   (c) presented, on arrival at the border inspection post, to the official veterinarian and may not be removed from the border inspection post until the checks in accordance with Directive 90/675/EEC have been carried out to the satisfaction of the official veterinarian and all the checks have been paid for.
6. Consignments transhipped via a port or airport elsewhere in the Community and which in accordance with Article 8.4 of Council Directive 90/675/EEC have not been subject to