

Department for Environment, Food and Rural Affairs

Notes for Guidance: Export Health Certificate for the entry into the European Union and Northern Ireland of breeding ratites and productive ratites 8438

June 2025

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No: 8438 NFG

EHC for entry into the EU or NI of breeding ratites and productive ratites.

NOTES FOR GUIDANCE (NFG) FOR THE CERTIFYING OFFICERS (CO) AND EXPORTERS

1. APPLICABLE LEGISLATION

[Decision 2007/240/EC](#)

[Regulation \(EU\) No 2016/429](#)

[Regulation \(EU\) No 2017/625](#)

[Regulation \(EU\) No 2018/1882](#)

[Commission Delegated Regulation \(EU\) 2019/2035](#)

[Commission Delegated Regulation \(EU\) 2020/689](#)

[Commission Delegated Regulation \(EU\) 2020/692](#)

[Commission Implementing Regulation \(EU\) 2020/2235](#)

[Commission Implementing Regulation \(EU\) 2021/404](#)

[Commission Implementing Regulation \(EU\) 2021/403](#)

[Regulation \(EU\) No 2018/1882](#)

[Implementing Regulation \(EU\) 2024/351 - Model EHC amending Implementing Regulation \(EU\) 2021/403](#)

[Commission Delegated Regulation \(EU\) 2023/905](#)

Any EU legislation referenced in the certificate must be complied with and EU legislation can be accessed on the following link. You should ensure you use the latest version: <https://eur-lex.europa.eu/homepage.html>

Please note that Official Control Regulations 2017/625 have repealed Regulation (EC) No 854/2004, 882/2004 and Directive No 96/23/EC. Please see link:

<https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32017R0625&from=EN>

Consolidated legislation

Consolidated texts, which integrate the basic instruments of EU legislation with their amendments and corrections in a single, non-official document, are available. Each consolidated text contains a list of all legal documents taken into account for its construction.

You can search for consolidated texts by using the 'find results by document number' option on the European Commission website. Once you have selected the relevant legislation, click 'document information', and then scroll down to 'all consolidated versions' and select the most recent version.

<https://eur-lex.europa.eu/homepage.html>

Please note that the consolidated text may not contain the latest amendment to the legislation, as it takes several weeks for this to be updated.

Texts provided in this section are intended for information only. Please note that these texts have no legal value. For legal purposes please refer to the texts published in the 'Official Journal of the EU'.

IMPORTANT

These notes provide guidance to COs and exporters. The NFG should have been issued to you together with the relevant export certificate applicable for dispatch of breeding or productive ratites to the EU or NI. The NFG should not be read as a standalone document but in conjunction with the veterinary certificate.

In addition to the relevant export certificate, ratites transiting through the EU or NI must be accompanied by certificate(s) required by the third country of destination. We strongly suggest that exporters obtain full details of the importing country's requirements from the veterinary authorities in the country concerned, or their representatives in the UK, in advance of each consignment.

[Please note, policies are being reviewed. NFG will be further amended to provide specific guidance. Traders should look at NFGs regularly for any updates]

2. SCOPE OF THE CERTIFICATE

This-certificate is for movements into the EU or NI of breeding or productive ratites.

It may also be used for ratites transiting the EU to another third country

The certifying OV must check the UK's status in the third country list in Annex 5 of Regulation (EU) 2021/404 as amended by (EU) 2021/634 against the commodity he/she is certifying and ensure that the relevant provisions are certified accordingly.

The conditions for import to the EU of breeding or productive ratites are different for consignments of less than 20 units, please refer to Article 4 of Regulation (EU) 2021/404 for further information.

This certificate is to be completed according to the notes for the completion of certificates provided for in Chapter 4 of Annex I to Implementing Regulation (EU) 2020/2235

3. CERTIFICATION BY AN OV

In **England, Scotland and Wales**, this certificate must be signed by a Government Veterinary Officer (e.g. APHA, FSA or FSS employed veterinary officers) or by an Official Veterinarian (OV) appointed by APHA on behalf of Ministers in Defra, the Scottish Government or the Welsh Government and who hold the appropriate Official Controls Qualification (Veterinary) (OCQ (V)) authorisation.

OVs must sign and stamp, with the OV stamp, the health certificate in ink of a different colour to that of the printing of the EHC. There is no requirement to sign and stamp in a specific colour.

The OV should keep a copy of the signed certificate and any supporting documents for at least two years after signature or receipt/dispatch of the consignment, whichever is later. These can be electronic copies.

EHC in foreign language/s of the EU Member States (MSs).

EHC should be in English and the foreign language of the Border Control Post (BCP) of entry in the EU. The original copy of the required EHC must accompany the consignment to the BCP of entry.

Listing of the EU MS BCPs can be found here: https://ec.europa.eu/food/animals/vet-border-control/bip-contacts_en

The foreign language certificate as received from the APHA Centre for International Trade at Carlisle or via the Export Health Certificates Online system (EHCO) and bearing the same unique reference number as the English certificate, should be considered an official and accurate translations of the English, as published in EU legislation.

The (sub-) paragraphs / options and how they are numbered and formatted is identical in the English and foreign language editions and to the legislation published by the European Commission. Therefore, the same phrases/sentences in the foreign language versions as in the English version should be struck through and these deletions should be stamped and initialled in both versions. Both versions must also be signed (as opposed to being initialled) and stamped by the OV, the foreign language certificate is deemed to be a genuine and properly authorised translation of the English version.

This also applies to any instructions in the guidance notes to strike out certain paragraphs or to certify statements that the country is free of certain notifiable diseases etc.

Additional information can be found in the APHA Vet Gateway:

http://apha.defra.gov.uk/External_OV_Instructions/Export_Instructions/Certification_Procedures/index.htm

SIGNING AND STAMPING

When signing a certificate, the CO should ensure that the certificate contains no deletions or alterations, other than those which are indicated on the certificate to be permissible and any corrections to permitted entries, subject to such changes being initialled and stamped (in the margin) by the CO. Permissible deletions are normally indicated in the 'Notes' section at the end of the certificate, with the instruction 'Keep as appropriate' or 'delete if not applicable'.

- Where the certificate contains optional or contextual statements, the statements which are not relevant shall be crossed out, individually initialled and stamped by the CO, or completely removed from the certificate.
- Permitted paragraphs and sections may be crossed out by applying a 'Z' across the section or paragraph rather than crossing out line by line.
- There is no requirement for a date and time to accompany each stamp. The date is only entered at the required entry field in Part I of the certificate, and at the end where the CO signs, stamps and dates that action.
- We are aware of some BCPs demanding that all handwritten information in Part 1 of the EHC is initialled and stamped, including handwritten scoring out of otherwise blank boxes. There is no legal requirement in EU legislation that all the hand-written information entered in the certificate must be signed and stamped. It is only in the case of correction, in any part of the certificate, or in the case of statements to be crossed out, that the certifier must add signature (or initials) and stamp. This has been confirmed by the European Commission. The Commission noted however, in the case of a hand-written certificate, it is expected that the same one person completes the document. If not, the BCP might suspect that empty boxes were completed by another person after the certificate has been signed by the official.

You should consider checking with the specific BCP regarding their preference when it comes to the stamping and initialling of handwritten scoring out of otherwise blank boxes in Part I of the EHC.

- **Clarification from the European Commission means that all pages (as opposed to sheets of paper) are signed and stamped once individually in place of fan stamping and in addition to any permitted alterations. There is no requirement to fan stamp.**
- COs are reminded to consult the NFG prior to the certification of each EHC. NFG will be updated with this new information in due course.

Further Information COs should make sure they are familiar with all relevant guidance and other documents relating to EHCs and that they discuss requirements with exporters in advance.

See <http://apha.defra.gov.uk/official-vets/Guidance/exports/ehc-online.htm>

You can also contact APHA's Centre for International Trade (CIT) on 03000 200 301.

PART I: DETAILS OF THE CONSIGNMENT

All boxes in Part I of the certificate must be completed. When a box is not applicable/optional, and not filled, please score it through.

Please use schedule to be attached to the certificate if there is not enough space to fill the information. See Section 'Addition of Schedules' below.

Please complete all the boxes in Part I of the certificate in accordance with the guidance laid down in Chapter 4 of Annex I to [Commission Implementing Regulation \(EU\) 2020/2235](#), Amended by [Implementing Regulation \(EU\) 2023/2744](#).

The Harmonised System (HS) Code is a commodity classification system used as a basis for customs tariffs and for international trade statistics.

It is the exporter's responsibility to ensure that the HS code is entered correctly and accurately reflects the product(s) being consigned.

Further information on HS Codes can be found online at:

<https://www.gov.uk/trade-tariff/sections> and <http://madb.europa.eu/madb/euTariffs.htm>

PART II: CERTIFICATION

II.1 Animal Health Attestation

The OV signing the export veterinary certificate must ensure that the animal health attestation set out in Part II of the veterinary certificate have been complied with.

They must ensure that they are aware of the relevant provisions of Regulation (EU) 2019/2035, regarding the approval details of premises, health status of the flock of origin and guarantees for Newcastle disease and Avian Influenza (AI).

II.1.1 (a) – Enter the territory code. GB is listed for all of the relevant commodities. The relevant regulations are [Implementing Regulations \(EU\) 2021/404](#) and [2021/405](#). These regulations have been amended by [Implementing Regulations 2021/634](#) and [2021/606](#), adding the GB and the Crown Dependencies to the relevant lists.

II.1.1(b) - This can be signed based on the compliance of the UK's surveillance programme with EU guidelines and recognised compliance with Regulation (EU) 2019/2035 requirements.

II.1.1(c) - [See Section 4](#). Notifiable Disease Clearance

II.1.2 – The first option may be certified based on the notifiable disease clearance in [Section 4](#).

II.1.3(a) - The first option may be signed. This can be signed on the basis that vaccination against AI is prohibited in GB. There is no plan to apply vaccination against AI in the UK even in an outbreak situation.

II.1.3(b) - The first option may be certified on the basis that the ratites have come from the UK. Annex XV of Regulation (EU) 2020/692 relates to the pathogenicity index of the

Newcastle disease vaccine. Any vaccine used in the flock of origin must have a marketing authorisation issued by the Veterinary Medicines Directorate (VMD) of DEFRA or the equivalent licensing body in another EU Member State and therefore must have an intracerebral pathogenicity index of less than 0.4. Where the ratites have been imported additional enquiries must be made to ensure that the vaccines have a suitably low pathogenicity index or that the additional statements can be satisfied.

II.1.4 – Requires that the flock of origin has been resident in the zone for a continuous period of at least 3 months immediately prior to the date of loading for dispatch to the Union or since hatching if they are less than 3 months of age. This may be certified after the OV has checked the movement records of the establishment. If animals have been introduced into the zone from **third countries**, the OV must have sight of the import permits to be satisfied that the health status of the imported flock meets those standards of Regulation (EU) 2016/429 and Regulation (EU) 2020/692 and that the country / zone of origin is listed in the Part 1. Section B of Annex V to [Implementing Regulation \(EU\) 2021/404](#). The OV would be advised to keep copies of the import permits for their records. If animals were introduced into the zone from the **EU Member State(s)**, this can be certified after the OV has checked movement records of the establishment and had sight of relevant Import Health Certificate(s), to establish the origin of the imported animals. The OV would be advised to keep copies of the documents for their records.

II.1.5 - The establishment must be named and supplied with an approval number that appears on a list of establishments drawn up and published by the Commission. This name and unique approval number must be in Box I.11 and in GB this refers to the Poultry Health Scheme (PHS) approval number.

II.1.5(a) - Check that the approval has not been suspended

II.1.5(b) & (c) - Should be certified on the basis that the farm and animal establishments are registered and approved by APHA and receive regular animal health visits from a farm veterinarian. Frequency of such visitation is proportionate to the risk. Records of animals should be kept for 3 years.

II.1.5(d) & (e) - May be certified on the basis of notifiable disease clearances, as above, as the birds came from holdings in the UK.

II.1.5(f) - See Section 4. Notifiable disease clearances for the most up to date health status of the zone.

II.1.6(a) - May be certified on the basis that vaccination for AI in the UK is not permitted. There is no plan to apply vaccination against AI in the UK even in an outbreak situation.

II.1.6(b) - May be certified after the OV has checked the establishments records. One option must be deleted. The OV is advised to keep a copy of the records to support their certification. If the flock has been vaccinated against Newcastle disease, then the second option must be certified, and the table completed.

II.1.6(c) - The clinical examination must be done within 24 hours of loading for dispatch to the EU. See Section 8 Clinical Examination. The OV must be familiar with the diseases listed in [Article 5 of Regulation \(EU\) 2016/429 and listed in Annex II](#) thereof, also the diseases as listed in [the Annex to Commission Implementing Regulation \(EU\) 2018/1882](#).

II.1.7 - May be certified as required on inspection of the establishment records and on receipt of an owner's declaration.

II.1.8 - May be certified as required on inspection of the establishment records and on receipt of an owner's declaration.

II.1.9 - May be certified on the basis of notifiable disease clearances, as above, as the animals came from holdings in the UK.

II.1.10 - Enter the date of the clinical inspection which should be within 24 hours of loading for dispatch to the EU. This may be certified if II.1.6(c) has been certified. The clinical examination must be done within 24 hours of loading and may only be performed by an OV of the country of origin. The OV should be vigilant for highly pathogen AI and Newcastle disease.

II.1.11 - May be certified if all of the points (a)-(e) are complied with at time of loading. Containers must be labelled with name and ISO code of the third country of origin, number of birds, name, address and approval number of the establishment of origin, specific identification number of the container, name of the Member State of destination.

II.1.12 - Enter the date of dispatch to the EU.

The certifying Official Veterinarian must ensure that the transport was cleaned and disinfected with an authorised disinfectant before loading in accordance with the relevant provisions of Assimilated EU Regulation No 1/2005 and that other parts of the attestation are complied with. See Section 7 on Animal Transport Attestation and gov.uk for further information on approved disinfectants. Every animal should be fit for the journey that is planned. A declaration from the owner / transporter must be sought to confirm relevant requirements have been met.

II.1.13 - This may be deleted. This section is only intended for the export of consignments of birds being exported to Member States granted the status free from Newcastle disease virus without vaccination in accordance with Article 66 of Commission Regulation (EU) 2020/689 and comply with the points listed. The certifying OV would need to seek the relevant evidence/access to records to satisfy themselves that these requirements on these points are complied with. (d) the tests should be carried out on samples taken by or under the control of the competent authority. Taking and testing of samples can be found; <http://apha.defra.gov.uk/vet-gateway/tte/nad.htm>

Further guidance for completion of this certificate can be found in Chapter 4 of Annex II to Commission Implementing Regulation (EU) 2021/404.

II.2 - This attestation on antimicrobial medicinal products is added which must be crossed out or deleted until 3 September 2026.

4. NOTIFIABLE DISEASE CLEARANCE

Commodities of poultry or poultry meat can be exported into the EU from the territory code listed in column 2 of the table in Part 1 of Annex V to [Commission Implementing Regulation \(EU\) 2021/404](http://Commission%20Implementing%20Regulation%20(EU)%2021/404). Ensure you are looking at the most up to date version of the Regulation. If the latest consolidated version does not include the latest amendment, this amendment needs to be looked at separately.

If the commodity to be exported is listed against GB-0, it can be exported to the EU from the whole territory of the UK. You will have to insert “GB-0” into the “territory code” box on the EHC.

If the commodity to be exported is listed against GB-1, it means that the UK is being regionalised because of a disease outbreak. All premises of origin (eg Flocks of origin, slaughter house, processing or storage premises as applicable) have to be located in GB-1. The OV must ensure that this information is correct. For up-to-date GB-1 and GB-2 areas please refer to the online interactive map where you have to check whether the premises of origin are all within the GB-1 area using the premises post codes. The interactive map can be found in the link below under “**COs Obtaining Clearance for AI**”.

<http://apha.defra.gov.uk/official-vets/Guidance/exports/ehc-online.htm>

Areas listed under GB-2 (and detailed as GB-2.1, GB-2.2 etc.) are restricted from exports between the “closing” and “opening” dates listed against those areas.

Some export certificates for animals and animal products will include statements that will require the OV to certify that specified areas or the entire country of origin are free from certain diseases.

Where it is possible for the CO (OV or Environmental Health Officer (EHO)) in Great Britain to obtain disease clearance themselves, the Centre for international Trade – Carlisle (CITC) will not issue a 618NDC notifiable disease clearance.

COs must check the following sources of disease information for the United Kingdom immediately prior to certification, to ensure disease freedom statements can be certified:

- the Notifiable Disease Occurrence List for Great Britain (ET171) available on the [Exports > Certification Procedures](#) page of the APHA Vet Gateway
- the UK Status for Non-Notifiable Diseases Relevant to Export Certification (ET152) available on the [Exports > Certification Procedures](#) page of the APHA Vet Gateway.
- For any postcodes in Northern Ireland, COs can obtain clearance using the interactive map provided by DAERA that can be found here: [AI Trade Map](#)

For Great Britain:

In the absence of a specific Notifiable Disease Clearance (618NDC) from CITC: COs may certify that the UK has disease free status or region free status for those diseases mentioned in the health certificate, once they have checked the disease list(s) for the last occurrence of the disease, and have ensured it complies with the time frames in the certificate.

In the event of a disease outbreak that affects a CO being able to obtain their own disease clearance, CITC will notify COs to make it clear which disease freedom statements should not be certified and where necessary, will issue a 618NDC notifiable disease clearance if the EHC can continue to be issued for certain regions that retain free status.

In the event of a disease outbreak after the EHC has been issued that affects the disease clearance, COs must not certify the EHC and must contact CITC immediately for advice on whether certification can still take place. If a disease outbreak affects the CO disease clearance procedures for this EHC, a 618NDC will be reinstated by CITC which will be issued with the EHC until a time when CO disease clearance can be reinstated.

NOTE: This does not apply to Transmissible Spongiform Encephalopathies (TSEs) or Bovine Tuberculosis (TB) freedom statements.

5. COLLECTION OF EVIDENCE

Certification Support Officers may not be utilised for gathering evidence relating to this certificate.

6. CONSIGNMENTS OR PARTS OF THE CONSIGNMENT ORIGINATING FROM NI, EU MEMBER STATES OR FROM A THIRD COUNTRY [WHEN APPLICABLE]

NI origin:

Consignments could potentially contain animals which have originated in NI. The certificate/documentation which the animal arrives into GB with may not contain sufficient information for the GB CO to sign the EU EHC.

Disease clearance for animals originating in NI can be completed using auto-clearance NDC found here:

<https://www.daera-ni.gov.uk/articles/notifiable-diseases-northern-ireland>

Where regional or local level disease clearance is required, this can be certified upon request on the basis of information from NI in the form of a declaration or a supporting health attestation.

Animal health statements which refer to the prohibition of certain vaccination programmes e.g. against FMD or CSF or ASF can be certified at a national level by the CO on the basis that NI also enforces a ban on such vaccinations in accordance with EU regulations.

Statements relating to implementation of a national system for identification and registration of livestock (cattle, sheep, goats, pigs, poultry) can be certified on the basis of the requirement to register all livestock animal births, moves and deaths on the DAERA database.

EU origin:

It is possible that some consignments may contain animals that are of EU origin and were imported into GB on a GB EHC. The GB EHC may not contain enough information to allow the CO to sign an EU EHC.

In such cases, the CO will need further information from the EU member state regarding particular attestations on the EHC that cannot be signed by the CO without support documentation. Thus, the GB exporter must request from the EU exporter an attestation or

written declaration from an EU registered vet. The GB exporter may wish to obtain these directly from the EU vet who has inspected the animals before export from the EU.

This supporting information must be in writing and kept by the GB CO. The GB CO is not required to attach it as a supporting document to the EHC, unless requested by the EU BCP or told otherwise.

Third country origin:

It is also possible that some consignments may contain animals that have been imported to GB from non-EU countries and fulfilled a residency period in GB, and GB exporters intend to export then to the EU. In these cases, COs may obtain a copy of the EHC for the import of such animals from the Third Country to GB.

GB COs are not required to attach a copy of the Third Country EHC as a supporting document to the EHC, unless requested by the EU BCP or specifically instructed in the NFG.

It is the GB exporter's ultimate responsibility to obtain any necessary support documents (from the EU member state exporter/Third Country exporter), to enable GB COs to be able to certify the live animals in good time before the export to the EU.

7. DECLARATION BY MASTER OF THE SHIP

A declaration by the master of the ship, as set out in Chapter 1 of Annex III of Regulation 2021/403, shall be attached to veterinary certificates for imports into the EU of ratites where the transport of those commodities includes transport by ship, even for part of the journey. You can find Master of the ship declaration here: <https://www.gov.uk/export-health-certificates/master-of-the-vessel-declaration-8466>

8. CLINICAL EXAMINATION

The inspection must be carried out within 24 hours of loading. The pre-export inspection should consist of a visual appraisal and, if deemed appropriate, physical examination of the animals for export. Each animal subject to an inspection must be assessed as an individual.

OVs must use their professional judgement to determine the level of inspection required in order to ensure that no animal is exported which shows signs of infectious disease and that animals are fit to travel to their intended destination.

9. ANIMAL TRANSPORT ATTESTATION

Council Regulation (EC) No 1/2005 and No 1/2002 is implemented under the

Welfare of Animals (Transport) (England) Order 2006 and parallel legislation in Scotland and Wales. If transported by air, animals should be transported in accordance with International Air Transport Association (IATA) standards. Every animal should be fit for the journey that is planned. Animals should be in good health, free of illness, free of significant wounds and able to walk without pain on all legs. Animals that are in sufficiently good health, should be able to withstand the stress of a journey without experiencing any unnecessary

pain or distress, and should arrive at their destination in good health. Animals that are injured (i.e. they are unable to move independently without pain or to walk unassisted) shall not be considered fit for transport.

10. ANIMAL HEALTH SCHEMES

Salmonella Control in Poultry

Regulation (EC) No 2160/2003 on the control of Salmonella in poultry is currently implemented through the UK Salmonella National Control Programme that is enforced by the Control of Salmonella in Poultry Order Regulation 2007 (England), the Control of Salmonella in Poultry (Wales) Order 2008, the Control of Salmonella in Poultry (Breeding, Laying and Broiler Flocks) (Scotland) Order 2009, the Control of Salmonella in Broiler Flocks Order 2009, and the Control of Salmonella in Turkey Flocks Order.

For consignments intended to be exported to Finland and Sweden, compliance with Commission Decision 2003/644 (EC) and Commission Decision 2004/235 must be certified. The OV must check the flock records to confirm that the appropriate tests have been carried out at the correct frequency with negative results of zoonotic salmonella species.

Concerning the results of testing, it should be described as positive ONLY if:

- In the case of breeding flocks, *S.hadar*, *S.virchow*, or *S.infantis* are detected.
- In the case of productive poultry, *S.enteritidis* or *S.typhimurium* are detected.

Poultry Health Scheme

Delegated Regulation (EU) 2019/2035 is currently being committed to through the equivalent Poultry Health Scheme (PHS) in Great Britain. A list of approved Poultry Health Scheme members can be found on the link below:

<https://www.gov.uk/government/publications/poultry-health-scheme-list-of-members>

Relevant text can be certified based on the establishment committing to the Poultry health scheme for the control and surveillance of specified non-zoonotic mycoplasma and salmonella bacterial species.

11. ADDITION OF SCHEDULES:

When the space in Part I or Part II of the certificate is insufficient to accommodate full details of the consignment a schedule may be used. In the relevant section of the certificate the CO should annotate the certificate 'see attached schedule'. A new schedule should be created (typed or clearly written) containing the same information as that required in the certificate. The schedule must include the certificate reference number on each page and must be signed, dated and stamped by the CO in a colour other than the printed text on each page and under the last entry. The schedule forms part of the certificate. All pages of the certificate, including the schedule, must be sequentially numbered. Any blank spaces in the schedule or the certificate should be struck through with diagonal lines.

Further guidance is available here:
http://apha.defra.gov.uk/External_OV_Instructions/Export_Instructions/Certification_Procedures/index.htm

12. CERTIFIED COPIES OF EHCs

When completing export certification, the CO and, if applicable, FCCO must make photocopies of, or scan and save all documents they certify. OV's must retain copies of certification documents in accordance with RCVS Certification principles.

<https://www.rcvs.org.uk/setting-standards/advice-and-guidance/code-of-professional-conduct-for-veterinary-surgeons/supporting-guidance/certification/>

COs must retain copies of all export documentation for a period of two years. A certified copy of this EHC does not need to be returned to the APHA CITC. For the purposes of completing routine Quality Assurance checks on export certification, CITC may request certified copies of certification from COs.

Further details on Post Certifying Procedures, 'certified copies' of certification and the types of documents that should be retained by COs can be found on the [APHA Vet Gateway](#).

13 LEGAL STATEMENT

The existing EU legislation that the UK complied with prior to the end of the Transition Period has been incorporated into our domestic law as "assimilated EU law" under the European Union (Withdrawal) Act 2018. References in our guidance and certification to such EU instruments should be taken to be references to this "assimilated EU law". The EU standards that this legislation includes continue to remain in force, without substantive amendment, as part of UK domestic law (apart from corrections to make the EU legislation fully operable

14. DISCLAIMER

This certificate and NFG are provided on the basis of information available at the time and may not necessarily comply fully with the requirements of the importing country. It is the exporter's responsibility to check the certificate against any relevant import permit or any advice provided by the competent authority in the importing country. If these do not match, the exporter should contact APHA in Carlisle.

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This publication is available at <https://www.gov.uk/search/all> Any enquiries regarding this publication should be sent to us at liveanimalexports.carlisle@apha.gov.uk

PB 8438 NFG

Version History

EHC

Published 30 August 2024

Part II:

II.2 - Attestation about the administration of antimicrobial medicinal products is added.

Notes - Footnote 13 is added.

Published 31 July 2024

II.1.4. – Updated to now allow the introduction of flocks to zone from a member state. Previously only from a third country.

NFG

Version 7: Published 16 June 2025

NOTIFIABLE DISEASE CLEARANCE – Section amended to include reference to AI map for NI.

Version 6: Published 30 August 2024

Applicable Legislation: Commission Delegated Regulation (EU) 2023/905 added

Part II: II.2 - Guidance is added about the attestation related to antimicrobial medicinal products.

Version 5: Published 31 July 2024

Applicable Legislation: Implementing Regulation (EU) 2024/351 added

Part 2 II.1.4 – Updated to reflect addition of member state flock introduction to zone