**Model animal health certificate for the entry into the Union of DOGS, CATS AND FERRETS (model ”CANIS-FELIS-FERRETS”)**

*Mudell taċ-ċertifikat tas-saħħa tal-annimali għad-dħul fl-Unjoni ta’ KLIEB, QTATES U INMSA (mudell “CANIS-FELIS-FERRETS”)*

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| **COUNTRY / PAJJIŻ:** Australia | **Animal health certificate to the EU / *Ċertifikat tas-saħħa tal-annimali lill-UE*** |
| **Part I: Description of consignment / *Parti I: Deskrizzjoni tal-kunsinna*** | **I.1** | **Consignor/Exporter / *Konsenjatur/Esportatur*** |   | **I.2** | **Certificate reference / *Referenza taċ-ċertifikat*** | **I.2a** | **IMSOC reference / *Referenza tal-IMSOC*** |
|  | Name / *Isem* |  |  |  |
|  | Address / *Indirizz* |  | **I.3** | **Central Competent Authority / *Awtorità Kompetenti Ċentrali***Department of Agriculture, Fisheries and Forestry |  | **QR CODE / *KODIĊI QR*** |
|  |  |
|  | Country / *Pajjiż*Australia | ISO country code / *Kodiċi ISO tal-pajjiż*AU | **I.4** | **Local Competent Authority / *Awtorità Kompetenti Lokali***Department of Agriculture, Fisheries and Forestry |  |  |
| **I.5** | **Consignee/Importer / *Konsenjatarju/Importatur*** |  | **I.6** | **Operator responsible for the consignment / *Operatur responsabbli għall-kunsinna*** |  |
|  | Name / *Isem* |  |  | Name / *Isem* |  |
|  | Address / *Indirizz* |  |  | Address / *Indirizz* |  |
|  | Country / *Pajjiż* | ISO country code / *Kodiċi ISO tal-pajjiż* |  | Country / *Pajjiż* | ISO country code / *Kodiċi ISO tal-pajjiż* |
| **I.7** | **Country of origin / *Pajjiż tal-oriġini***Australia | ISO country code / *Kodiċi ISO tal-pajjiż*AU | **I.9** | **Country of destination / *Pajjiż tad-destinazzjoni*** | ISO country code / *Kodiċi ISO tal-pajjiż* |
| **I.8** | **Region of origin / *Reġjun tal-oriġini***Australia | Code / *Kodiċi*AU-0 | **I.10** | **Region of destination / *Reġjun tad-destinazzjoni*** | Code / *Kodiċi* |
| **I.11** | **Place of dispatch / *Post tad-dispaċċ*** |  | **I.12** | **Place of destination / *Post tad-destinazzjoni*** |  |
|  | Name / *Isem* | Registration/Approval No / *Nru tar-Reġistrazzjoni/tal-Approvazzjoni* |  | Name / *Isem* | Registration/Approval No / *Nru tar-Reġistrazzjoni/tal-Approvazzjoni* |
|  | Address / *Indirizz* |  |  | Address / *Indirizz* |  |
|  | Country / *Pajjiż*Australia | ISO country code / *Kodiċi ISO tal-pajjiż*AU |  | Country / *Pajjiż* | ISO country code / *Kodiċi ISO tal-pajjiż* |
| **I.13** | **Place of loading / *Post tat-tagħbija*** | **I.14** | **Date and time of departure / *Data u ħin tat-tluq*** |
| **I.15** | **Means of transport / *Mezz tat-trasport*** |  |  | **I.16** | **Entry Border Control Post / *Post ta’ Kontroll fuq il-Fruntiera tad-Dħul*** |
|  | Aircraft / *Inġenju tal-ajru* | Vessel / *Bastiment* | **I.17** | **Accompanying documents / *Dokumenti ta’ akkumpanjament*** |
|  |  |
|  | Railway / *Ferrovija* | Road vehicle / *Vettura tat-triq* |  | Type / *Tip* | Code / *Kodiċi* |
|  | Identification / *Identifikazzjoni* |  | Country / *Pajjiż* | ISO country code / *Kodiċi ISO tal-pajjiż* |
|  | Commercial document reference / *Referenza tad-dokument kummerċjali* |  |
| **I.18** | **Transport conditions / *Kundizzjonijiet tat-trasport*** | Ambient / *Ambjentali* | Chilled / *Imkessaħ* | Frozen / *Iffriżat* |
| **I.19** | **Container number/Seal number / *Numru tal-kontenitur/Numru tas-siġill*** |
|  | Container No / *Nru tal-kontenitur* | Seal No / *Nru tas-siġill* |  |
| **I.20** | **Certified as or for / *Iċċertifikat bħala jew għal*** |
|  | Further keeping / *Żamma ulterjuri* |  |  |  |
|  |  | Confined establishment / *Stabbiliment konfinat* |  |  |
|  |  | Quarantine establishment / *Stabbiliment ta’ kwarantina* |  |  |
|  |  |  | Other / *Oħrajn* |  |
| **I.21** |  **For transit / *Għat-tranżitu*** | **I.22** |  **For internal market / *Għas-suq intern*** |
|  | Third country / *Pajjiż terz* | ISO country code / *Kodiċi ISO tal-pajjiż* | **I.23** |  |

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| **I.24** | **Total number of packages / *Għadd totali ta’ imballaġġi*** | **I.25** | **Total quantity / *Kwantità totali*** | **I.26** | **Total net weight/gross weight (kg) / *Piż nett totali/piż gross (kg)*** |
| **I.27** | **Description of consignment / *Deskrizzjoni tal-kunsinna*** |
| CN code / *Kodiċi CN* | Species / *Speċi* | Subspecies/Category / *Sottospeċi/Kategorija* | Sex / *Sess* | Identification system / *Sistema ta’ identifikazzjoni* | Identification number / *Numru ta’ identifikazzjoni* | Age / *Età* | Quantity / *Kwantità* |
|  |  |  |  |  |  |  |  |
|  |  |  |  | Nature of commodity / *Natura tal-komodità* |  |  |  |
|  |  |  |  |  |  | Test / *Test* |  |

| **COUNTRY / *PAJJIŻ***: Australia | **Certificate model CANIS-FELIS-FERRETS / *Mudell taċ-ċertifikat CANIS-FELIS-FERRETS*** |
| --- | --- |
| **Part II: Certification / *Parti II: Ċertifikazzjoni*** | **II. Health information / *Informazzjoni dwar is-Saħħa*** | **II.a**  | **Certificate reference / *Referenza taċ-ċertifikat*** | **II.b**  | **IMSOC reference / *Referenza tal-IMSOC*** |
| I, the undersigned official veterinarian hereby certify that the animals of the consignment described in Part I / *Jiena, il-veterinarju uffiċjali sottoskritt, b’dan qed niċċertifika, li l-annimali tal-kunsinna deskritta fil-Parti I*:II.1. come from a third country or territory, zone thereof with code: AU - 0(1) which, on the date of issue of this animal health certificate is authorised for the entry into the Union of dogs, cats and ferrets and is listed in Part 1 of Annex VIII to Commission Implementing Regulation (EU) 2021/404;*II.1. ġejjin minn pajjiż terz jew territorju, jew żona tiegħu bil-kodiċi: AU - 0(1) li, fid-data tal-ħruġ ta’ dan iċ-ċertifikat tas-saħħa tal-annimali hija awtorizzata għad-dħul fl-Unjoni ta’ klieb, qtates u inmsa u hija elenkata fil-Parti 1 tal-Anness VIII tar-Regolament ta’ Implimentazzjoni tal-Kummissjoni (UE) 2021/404;*(2) *either* [II.2. have been dispatched to the Union directly from the establishment of origin without passing through any other establishment;]*(2) jew [II.2. ikunu ġew dispaċċati lejn l-Unjoni direttament mill-istabbiliment tal-oriġini mingħajr ma jkunu għaddew minn xi stabbiliment ieħor;]*(2)(3) *or* [II.2. have undergone one single assembly operation in the country or territory, or zone thereof of origin which took place for not more than 6 days in an establishment fulfilling the following requirements:* it is approved for conducting assembly operations of dogs, cats and ferrets by the competent authority in the third country or territory in accordance with Article 10 of Commission Delegated Regulation (EU) 2019/2035;
* it has a unique approval number assigned by the competent authority of the third country or territory;
* it is listed for that purpose by the competent authority of the third country or territory of dispatch to the Union, including the information set out in Article 21 of Delegated Regulation (EU) 2019/2035;
* it complies with the record keeping requirements provided for in Article 73(2), point (a)(iv), of Delegated Regulation (EU) 2020/692;]

*(2)(3) jew [II.2. għaddew minn operazzjoni waħda ta’ assemblaġġ fil-pajjiż jew territorju, jew żona tiegħu tal-oriġini li seħħet għal mhux aktar minn 6 ijiem fi stabbiliment li jissodisfa r-rekwiżiti li ġejjin:** *huwa approvat għat-twettiq tal-operazzjonijiet ta’ assemblaġġ ta’ klieb, qtates u inmsa mill-awtorità kompetenti fil-pajjiż terz jew territorju f’konformità mal-Artikolu 10 tar-Regolament Delegat tal-Kummissjoni (UE) 2019/2035;*
* *għandu numru tal-approvazzjoni uniku assenjat mill-awtorità kompetenti tal-pajjiż terz jew territorju;*
* *huwa elenkat għal dak l-iskop mill-awtorità kompetenti tal-pajjiż terz jew territorju tad-dispaċċ lejn l-Unjoni, inkluża l-informazzjoni stabbilita fl-Artikolu 21 tar-Regolament Delegat (UE) 2019/2035;*
* *jikkonforma mar-rekwiżiti għaż-żamma ta’ rekords previsti fl-Artikolu 73(2), il-punt (a)(iv) tar-Regolament Delegat (UE) 2020/692;]*

(2)(3) *or* [II.2. have been dispatched from an animal shelter fulfilling the following requirements: * it is approved by the competent authority in the third country or territory in accordance with Article 11 of Delegated Regulation (EU) 2019/2035;
* it has a unique approval number assigned by the competent authority of the third country or territory;
* it is listed for that purpose by the competent authority of the third country or territory of dispatch, including the information provided for in Article 21 of Delegated Regulation (EU) 2019/2035;]

*(2)(3) jew [II.2. ġew dispaċċati minn santwarju tal-annimali li jissodisfa r-rekwiżiti li ġejjin:* * *huwa approvat mill-awtorità kompetenti fil-pajjiż terz jew territorju f’konformità mal-Artikolu 11 tar-Regolament Delegat (UE) 2019/2035;*
* *għandu numru tal-approvazzjoni uniku assenjat mill-awtorità kompetenti tal-pajjiż terz jew territorju;*
* *huwa elenkat għal dak l-iskop mill-awtorità kompetenti tal-pajjiż terz jew territorju tad-dispaċċ inkluża l-informazzjoni stabbilita fl-Artikolu 21 tar-Regolament Delegat (UE) 2019/2035;]*

(3) [II.3. have been loaded for dispatch to the Union on \_\_\_/\_\_\_/\_\_\_\_(dd/mm/yyyy) (4) in a means of transport which was cleaned and disinfected prior to loading with a disinfectant authorised by the competent authority in the third country or territory and constructed in such a way that:* animals cannot escape or fall out;
* visual inspection of the space where animals are kept is possible;
* the escape of animal excrements, litter or feed is prevented or minimized;]

*(3) [II.3. ikunu tgħabbew għad-dispaċċ lejn l-Unjoni fi \_\_\_/\_\_\_/\_\_\_\_ (jj/xx/ssss) (4) f’mezz ta’ trasport li jkun tnaddaf u ġie ddiżinfettat qabel it-tagħbija b’diżinfettant awtorizzat mill-awtorità kompetenti fil-pajjiż terz jew territorju u li jkun inbena b’tali mod li:** *l-annimali ma jkunux jistgħu jaħarbu jew jaqgħu;*
* *ikun possibbli li ssir spezzjoni viżwali tal-ispazju fejn jinżammu l-annimali;*
* *jiġi evitat jew minimizzat il-ħruġ ta’ eskrementi, mifrex jew għalf tal-annimali;]*

II.4 have been subjected with negative result to a clinical inspection, carried out by an official veterinarian in the third country or territory, or zone thereof of origin within the last 48 hours prior to the time of loading for dispatch to the Union for the detection of signs indicative of the occurrence of diseases, including the listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species and emerging diseases;*II.4 ikunu ġew soġġetti b’riżultat negattiv għal spezzjoni klinika, imwettqa minn veterinarju uffiċjali fil-pajjiż terz jew territorju jew żona tiegħu tal-oriġini fl-aħħar 48 siegħa qabel il-ħin tat-tagħbija għad-dispaċċ lejn l-Unjoni għad-detezzjoni ta’ sinjali li jindikaw l-okkorrenza ta’ mard, inklużi l-mard elenkat imsemmi fl-Anness I tar-Regolament Delegat tal-Kummissjoni (UE) 2020/692 rilevanti għall-ispeċi u l-mard emerġenti;*(2)*either* [II.5. are destined for direct entry into the Member State of destination to be isolated in:(2) *either* [a confined establishment;]](2) *or* [an approved quarantine establishment;]]*(2) jew [II.5. ikunu destinati għal dħul dirett fl-Istat Membru tad-destinazzjoni biex ikunu iżolati fi:**(2) jew [stabbiliment konfinat;]]**(2) jew [stabbiliment ta’ kwarantina approvat;]]*(2 )*or* [II.5. were at least 12 weeks old at the date of vaccination against rabies and at least 21 days have elapsed since the date of completion of the primary anti-rabies vaccination (5) carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council, and any subsequent revaccination was carried out within the period of validity of the preceding vaccination (6), and:*(2 )jew [II.5. kellhom mill-inqas 12-il ġimgħa fid-data tat-tilqim kontra r-rabja u jkunu għaddew mill-inqas 21 jum mid-data tat-tlestija tat-tilqim primarju kontra r-rabja(5) mwettaq f’konformità mar-rekwiżiti ta’ validità stabbiliti fl-Anness III tar-Regolament (UE) 576/2013 tal-Parlament Ewropew u tal-Kunsill, u kwalunkwe tilqim mill-ġdid sussegwenti twettaq fil-perjodu ta’ validità tat-tilqim preċedenti(6), u:*(2)*either* [come from, and in the case of transit are scheduled to transit through, a third country or territory listed in Annex II to Commission Implementing Regulation (EU) No 577/2013 and details of the relevant anti-rabies vaccination(s) are provided in columns 1 to 7 in the table below;]]*(2) jew [ikunu ġejjin minn, u f’każ ta’ tranżitu, ikunu skedati għal tranżitu minn, pajjiż terz jew territorju elenkat fl-Anness II tar-Regolament ta’ Implimentazzjoni tal-Kummissjoni (UE) Nru 577/2013 u d-dettalji tat-tilqim attwali kontra r-rabja huma pprovduti fil-kolonni minn 1 sa 7 fit-tabella ta’ hawn taħt;]]*(2)*or* [come from or are scheduled to transit through a third country or territory not listed in Annex II to Commission Implementing Regulation (EU) No 577/2013, and: *(2) jew [ikunu ġejjin minn jew ikunu skedati għal tranżitu minn, pajjiż terz jew territorju mhux elenkat fl-Anness II tar-Regolament ta’ Implimentazzjoni tal-Kummissjoni (UE) Nru 577/2013, u:* (a) the details of the relevant anti-rabies vaccination(s) are provided in columns 1 to 7 in the table below / *id-dettalji tat-tilqim rilevanti kontra r-rabja huma pprovduti fil-kolonni minn 1 sa 7 fit-tabella ta’ hawn taħt*, (b) a rabies antibody titration test (7), carried out on a blood sample taken by the veterinarian authorised by the competent authority not less than 30 days after the date of the preceding vaccination and at least 3 months prior to the date of issue of this animal health certificate, proved an antibody titre equal to or greater than 0,5 IU/ml (8) and any subsequent revaccination was carried out within the period of validity of the preceding vaccination, and the date of sampling for testing the immune response are provided in column 8 in the table below:]] / *test tat-titrazzjoni tal-antikorpi tar-rabja(7), imwettaq fuq kampjun tad-demm meħud mill-veterinarju awtorizzat mill-awtorità kompetenti mhux inqas minn 30 jum wara d-data tat-tilqim preċedenti u mill-inqas 3 xhur qabel id-data tal-ħruġ ta’ dan iċ-ċertifikat tas-saħħa tal-annimali, wera titru tal-antikorpi ta’ 0,5 IU/ml(8) jew akbar, u kwalunkwe tilqim mill-ġdid sussegwenti twettaq fil-perjodu ta’ validità tat-tilqim preċedenti, u d-data tat-teħid tal-kampjuni għall-ittestjar tar-reazzjoni immunoloġika hija pprovduta fil-kolonna 8 fit-tabella ta’ hawn taħt:]]*

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| --- | --- | --- | --- | --- | --- |
| **Transponder / *Transponder*** | **Date of vaccination [dd/mm/yyyy] / *Data tat-tilqim [jj/xx/ssss]*** | **Name and manufacturer of vaccine / *Isem u manifattur tal-vaċċin*** | **Batch number / *Numru tal-lott*** | **Validity of vaccination / *Validità tat-tilqima*** | **Date of blood sampling [dd/mm/yyyy] / *Data tat-teħid tal-kampjuni tad-demm [jj/xx/ssss]*** |
| **Alphanumeric code of the animal / *Kodiċi alfanumeriku tal-annimal*** | **Date of implantation and/or reading (9) [dd/mm/yyyy] / *Data tal-impjantazzjoni u/jew tal-qari(9) [jj/xx/ssss]*** |
| **From [dd/mm/yyyy] / *Minn [jj/xx/ssss]*** | **To [dd/mm/yyyy] / *Sa [jj/xx/ssss]*** |
| 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |
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(2)*either* [II.6. include dogs destined for a Member State listed in the Annex to Commission Implementing Regulation (EU) 2018/878 and those dogs have been treated against infestation with *Echinococcus multilocularis*, and the details of the treatment carried out by the administering veterinarian in accordance with point 2 of Annex XXI to Delegated Regulation (EU) 2020/692 (10) (11) are provided in the table below:*(2) jew [II.6. jinkludu l-klieb destinati għal Stat Membru elenkat fl-Anness tar-Regolament ta’ Implimentazzjoni tal-Kummissjoni (UE) 2018/878 u dawk il-klieb ġew trattati kontra l-infestazzjoni bl-Echinococcus multilocularis, u d-dettalji tat-trattament imwettaq mill-veterinarju li jamministra f’konformità mal-punt 2 tal-Anness XXI tar-Regolament Delegat (UE) 2020/692 (10) (11) huma pprovduti fit-tabella ta’ hawn taħt:*

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| **Transponder or tattoo. Alphanumeric code of the dog / *Transponder jew tatwaġġ. Kodiċi alfanumeriku tal-kelb*** | **Anti-Echinococcus treatment / *Trattament kontra l-echinococcus*** | **Administering veterinarian / *Veterinarju li jamministra*** |
| **Name and manufacturer of the product / *Isem il-prodott u manifattur tiegħu*** | **Date [dd/mm/yyyy] and time of treatment [00:00] / *Data [jj/xx/ssss] u l-ħin [00:00] tat-trattament*** | **Name in capitals, stamp and signature / *Isem b’ittri kapitali, timbru u firma*** |
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(2)*or* [II.6. include dogs which have not been treated against infestation with *Echinococcus multilocularis.*]*(2) jew [II.6. jinkludu l-klieb li ma ġewx trattati kontra l-infestazzjoni bil-Echinococcus multilocularis.]*(2)*or* [II.6. include dogs destined for direct entry into the Member State of destination to be isolated in:*(2) jew [II.6. jinkludu l-klieb destinati għad-dħul dirett fl-Istat Membru tad-destinazzjoni biex ikunu iżolati fi:*(1) *either* [a confined establishment.]]*(1) jew [stabbiliment konfinat.]]*(1) *or* [an approved quarantine establishment.]]*(1) jew [stabbiliment ta’ kwarantina approvat.]]***Notes:**This animal health certificate is intended for commercial entries into the Union of dogs, cats and ferrets, including when they are destined to a confined establishment or to an approved quarantine establishment and when the Union is not the final destination of the animals and for the entry into the Union of dogs, cats and ferrets moved in accordance with Article 5(4) of Regulation (EU) No 576/2013 of the European Parliament and of the Council.In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.This animal health certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 4 of Annex I to Commission Implementing Regulation (EU) 2020/2235.***Noti:****Dan iċ-ċertifikat tas-saħħa tal-annimali huwa maħsub għad-dħul kummerċjali fl-Unjoni ta’ klieb, qtates u inmsa, inkluż meta jkunu destinati għal stabbiliment konfinat jew għal stabbiliment ta’ kwarantina approvat u meta l-Unjoni ma tkunx id-destinazzjoni finali tal-annimali u għad-dħul fl-Unjoni ta’ klieb, qtates u inmsa mċaqilqa f’konformità mal-Artikolu 5(4) tar-Regolament (UE) Nru 576/2013 tal-Parlament Ewropew u tal-Kunsill.**F’konformità mal-Ftehim dwar il-ħruġ tar-Renju Unit tal-Gran Brittanja u l-Irlanda ta’ Fuq mill-Unjoni Ewropea u mill-Komunità Ewropea tal-Enerġija Atomika, u b’mod partikolari l-Artikolu 5(4) tal-Protokoll dwar l-Irlanda/l-Irlanda ta’ Fuq flimkien mal-Anness 2 ta’ dak il-Protokoll, ir-referenzi għall-Unjoni f’dan iċ-ċertifikat tas-saħħa tal-annimali jinkludu lir-Renju Unit fir-rigward tal-Irlanda ta’ Fuq.**Dan iċ-ċertifikat tas-saħħa tal-annimali għandu jimtela f’konformità man-noti għall-mili taċ-ċertifikati previsti fil-Kapitolu 4 tal-Anness I tar-Regolament ta’ Implimentazzjoni tal-Kummissjoni (UE) 2020/2235.***Part I:**Box reference I.20: Certified as or for: Indicate: * "Further keeping" where dogs, cats or ferrets are moved in accordance with Title V of Part II of Delegated Regulation (EU) 2020/692;
* Confined establishment: as defined in Article 4(48) of Regulation (EU) 2016/429 of the European Parliament and of the Council;
* Approved quarantine establishment: as defined in Article 3(9) of Commission Delegated Regulation (EU) 2020/688;
* "others" where dogs (*Canis lupus familiaris*), cats (*Felis silvestris catus*) or ferrets (*Mustela putorius furo*) are moved in accordance with Article 5(4) of Regulation (EU) No 576/2013 of the European Parliament and of the Council.

***Parti I:****Kaxxa ta’ referenza I.20: Iċċertifikat bħala jew għal: Indika:* * *“Żamma ulterjuri” fejn il-klieb, il-qtates jew l-inmsa jiġu mċaqilqa f’konformità mat-Titolu V tal-Parti II tar-Regolament Delegat (UE) 2020/692;*
* *Stabbiliment konfinat: kif definit fl-Artikolu 4(48) tar-Regolament (UE) 2016/429 tal-Parlament Ewropew u tal-Kunsill;*
* *Stabbiliment ta’ kwarantina approvat: kif definit fl-Artikolu 3(9) tar-Regolament Delegat tal-Kummissjoni (UE) 2020/688;*
* *“oħrajn”, fejn il-klieb (Canis lupus familiaris), il-qtates (Felis silvestris catus) jew l-inmsa (Mustela putorius furo) jiġu mċaqilqa f’konformità mal-Artikolu 5(4) tar-Regolament (UE) Nru 576/2013 tal-Parlament Ewropew u tal-Kunsill.*

**Part II:**(1) Code of the zone as it appears in column 2 of the table in Part 1 of Annex VIII to Implementing Regulation (EU) 2021/404.(2) Delete if not applicable.(3) Not applicable to the movement of dogs, cats and ferrets other than non-commercial movements, kept as pet animals in households that may not be carried out in accordance with the conditions laid down in Article 245(2) or Articles 246(1) and (2) of Regulation (EU) 2016/429.(4) Date of loading: it may not be a date prior to the date of authorisation of the zone for the entry into the Union, or a date in a period when restriction measures have been adopted by the Union against the entries into the Union of those animals from that zone.(5) Any revaccination shall be considered a primary vaccination if it was not carried out within the period of validity of a previous vaccination.(6) A certified copy of the identification and vaccination details of the animals concerned shall be attached to the animal health certificate.(7) The rabies antibody titration test referred to in point II.5:- shall be carried out on a sample collected by a veterinarian authorised by the competent authority, at least 30 days after the date of vaccination and 3 months prior to the date of dispatch to the Union*;*- shall measure a level of neutralising antibody to rabies virus in serum equal to or greater than 0,5 IU/ml;- shall be performed by an official laboratory;- shall not be renewed on an animal, which following that test with satisfactory results, has been revaccinated against rabies within the period of validity of a previous vaccination. A certified copy of the official report from the official laboratory on the result of the rabies antibody test referred to in point II.5 shall be attached to the animal health certificate.(8) By certifying this result, the official veterinarian confirms that he has verified, to the best of his ability and where necessary with contacts with the laboratory indicated in the report, the authenticity of the laboratory report on the results of the antibody titration test referred to in point II.5.(9) In conjunction with note (6), the marking of the animals concerned by the implantation of a transponder shall be verified before any entry is made in this animal health certificate and shall always precede any vaccination, or where applicable, testing carried out on those animals.(10) The treatment against infestation with *Echinococcus multilocularis* referred to in point II.6 shall:- be administered by a veterinarian within not more than 48 hours and not less than 24 hours prior to the time of the scheduled dispatch of the dogs to one of the Member States or parts thereof listed in the Annex to Commission Implementing Regulation (EU) 2018/878;- consist of an approved medicinal product which contains the appropriate dose of praziquantel or pharmacologically active substances, which alone or in combination, have been proven to reduce the burden of mature and immature intestinal forms of *Echinococcus multilocularis* in the host species concerned.(11) The table referred to in point II.6 shall be used to document the details of a further treatment if administered after the date the animal health certificate was signed and prior to the scheduled entry into one of the Member States or parts thereof listed in the Annex to Implementing Regulation (EU) 2018/878.***Parti II:****(1) Il-kodiċi taż-żona kif jidher fil-kolonna 2 tat-tabella fil-Parti 1 tal-Anness VIII tar-Regolament ta’ Implimentazzjoni (UE) 2021/404.**(2) Ħassar jekk mhux applikabbli.**(3) Mhux applikabbli għall-moviment ta’ klieb, qtates u inmsa għajr il-movimenti mhux kummerċjali, miżmuma bħala annimali domestiċi fl-unitajiet domestiċi li jista’ ma jitwettaqx f’konformità mal-kundizzjonijiet stabbiliti fl-Artikolu 245(2) jew fl-Artikoli 246(1) u (2) tar-Regolament (UE) 2016/429.**(4) Id-data tat-tagħbija: ma tistax tkun data qabel id-data tal-awtorizzazzjoni taż-żona għad-dħul fl-Unjoni, jew data f’perjodu meta jkunu ġew adottati miżuri ta’ restrizzjoni mill-Unjoni kontra d-dħul fl-Unjoni ta’ dawk l-annimali minn dik iż-żona.**(5) Kwalunkwe tilqim mill-ġdid għandu jitqies bħala tilqim primarju jekk ma jkunx twettaq fil-perjodu ta’ validità tat-tilqim preċedenti.**(6) Għandha tiġi mehmuża kopja ċċertifikata tad-dettalji ta’ identifikazzjoni u tilqim tal-annimali kkonċernati maċ-ċertifikat tas-saħħa tal-annimali.**(7) It-test tat-titrazzjoni tal-antikorpi tar-rabja msemmi fil-punt II.5:**- għandu jitwettaq fuq kampjun meħud minn veterinarju awtorizzat mill-awtorità kompetenti, mill-inqas 30 jum wara d-data tat-tilqim u 3 xhur qabel id-data tad-dispaċċ lejn l-Unjoni;**- għandu jkejjel livell ta’ antikorp newtralizzanti għall-virus tar-rabja fis-serum ta’ 0,5 IU/ml jew akbar;**- għandu jitwettaq minn laboratorju uffiċjali;**- ma għandux jerġa’ jitwettaq fuq annimal, li wara dak it-test b’riżultati sodisfaċenti, li jkun tlaqqam mill-ġdid kontra r-rabja fil-perjodu tal-validità ta’ tilqim preċedenti.* *Għandha tiġi mehmuża kopja ċċertifikata tar-rapport uffiċjali mil-laboratorju uffiċjali dwar ir-riżultat tat-test tal-antikorpi tar-rabja msemmi fil-punt II.5 maċ-ċertifikat tas-saħħa tal-annimali.**(8) Billi jiċċertifika dan ir-riżultat, il-veterinarju uffiċjali jikkonferma li huwa vverifika, bl-aħjar kapaċità tiegħu u, fejn meħtieġ, b’kuntatti mal-laboratorju indikat fir-rapport, l-awtentiċità tar-rapport tal-laboratorju dwar ir-riżultati tat-test tat-titrazzjoni tal-antikorpi msemmi fil-punt II.5.**(9) Flimkien man-nota (6), l-immarkar tal-annimali kkonċernati permezz tal-impjantazzjoni ta’ transponder għandu jiġi vverifikat qabel ma ssir kwalunkwe entrata f’dan iċ-ċertifikat tas-saħħa tal-annimali u dejjem għandu jsir qabel kwalunkwe tilqim, jew fejn applikabbli, ittestjar imwettaq fuq dawk l-annimali.**(10) It-trattament kontra l-infestazzjoni bl-Echinococcus multilocularis imsemmi fil-punt II.6 għandu:**- jiġi amministrat minn veterinarju f’mhux aktar minn 48 siegħa u mhux inqas minn 24 siegħa qabel il-ħin tad-dispaċċ skedat tal-klieb lejn wieħed mill-Istati Membri jew partijiet tagħhom elenkat fl-Anness tar-Regolament ta’ Implimentazzjoni tal-Kummissjoni (UE) 2018/878;**- ikun jikkonsisti fi prodott mediċinali approvat li jkun fih id-doża xierqa ta’ prażikwantel jew sustanzi farmakoloġikament attivi, li weħidhom jew flimkien, urew li jnaqqsu l-piż ta’ forom intestinali maturi u immaturi ta’ Echinococcus multilocularis fl-ispeċi ospitanti kkonċernata.**(11) It-tabella msemmija fil-punt II.6. għandha tintuża biex jiġu ddokumentati d-dettalji ta’ trattament ulterjuri jekk jiġi amministrat wara d-data li fiha ġie ffirmat iċ-ċertifikat tas-saħħa tal-annimali u qabel id-dħul skedat f’wieħed mill-Istati Membri jew partijiet tiegħu elenkat fl-Anness tar-Regolament ta’ Implimentazzjoni (UE) 2018/878.* |
| **Official veterinarian / *Veterinarju uffiċjali*** |
| Name (in capital letters) / *Isem (b’ittri kapitali)* |  |  |  |
| Date / *Data* |  | Qualification and title / *Kwalifika u titolu* |  |
| Stamp / *Timbru* |  | Signature / *Firma* |  |