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

*Deimhniú sláinte ainmhithe caighdeánach maidir le teacht isteach san Aontas MADRAÍ, CAT AGUS FIRÉAD (leagan caighdeánach ‘CANIS-FELIS-FERRETS’)*

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| **COUNTRY / *TÍR:*** Australia | **Animal health certificate to the EU / *Deimhniú sláinte ainmhithe don Aontas Eorpach*** |
| **Part I: Description of consignment / *Cuid I: Tuairisc ar an gcoinsíneacht*** | **I.1** | **Consignor/Exporter / *Coinsíneoir/Onnmhaireoir*** |   | **I.2** | **Certificate reference / *Tagairt an deimhnithe*** | **I.2a** | **IMSOC reference / *Tagairt CBFRO*** |
|  | Name / *Ainm* |  |  |  |
|  | Address / *Seoladh* |  | **I.3** | **Central Competent Authority / *Údarás Inniúil Lárnach***Department of Agriculture, Fisheries and Forestry |  | **QR CODE / *CÓD QR*** |
|  |  |
|  | Country / *Tír*Australia | ISO country code / *Cód tíre ISO*AU | **I.4** | **Local Competent Authority / *Údarás Inniúil Áitiúil***Department of Agriculture, Fisheries and Forestry  |  |  |
| **I.5** | **Consignee/Importer / *Coinsíní/Allmhaireoir*** |  | **I.6** | **Operator responsible for the consignment / *An t‑oibreoir atá freagrach as an gcoinsíneacht*** |  |
|  | Name / *Ainm* |  |  | Name / *Ainm* |  |
|  | Address / *Seoladh* |  |  | Address / *Seoladh* |  |
|  | Country / *Tír* | ISO country code / *Cód tíre ISO* |  | Country / *Tír* | ISO country code / *Cód tíre ISO* |
| **I.7** | **Country of origin / *An tír thionscnaimh***Australia | ISO country code / *Cód tíre ISO*AU | **I.9** | **Country of destination / *An tír cinn scríbe*** | ISO country code / *Cód tíre ISO* |
| **I.8** | **Region of origin / *An réigiún tionscnaimh***Australia | Code / *Cód*AU-0 | **I.10** | **Region of destination / *An réigiún cinn scríbe*** | Code / *Cód* |
| **I.11** | **Place of dispatch / *Áit seolta*** |  | **I.12** | **Place of destination / *Ceann scríbe*** |  |
|  | Name / *Ainm* | Registration/Approval No / *Uimh. Chlárúcháin/Formheasa* |  | Name / *Ainm* | Registration/Approval No / *Uimh. Chlárúcháin/Formheasa* |
|  | Address / *Seoladh* |  |  | Address / *Seoladh* |  |
|  | Country / *Tír*Australia | ISO country code / *Cód tíre ISO*AU |  | Country / *Tír* | ISO country code / *Cód tíre ISO* |
| **I.13** | **Place of loading / *Ionad luchtaithe*** | **I.14** | **Date and time of departure / *Dáta agus am imeachta*** |
| **I.15** | **Means of transport / *Modh iompair*** |  |  | **I.16** | **Entry Border Control Post / *Post Rialaithe Teorann Iontrála*** |
|  | Aircraft / *Aerárthach* | Vessel / *Soitheach* | **I.17** | **Accompanying documents / *Doiciméid tionlacain*** |
|  |  |
|  | Railway / *Iarnród* | Road vehicle / *Feithicil bhóthair* |  | Type / *Saghas* | Code / *Cód* |
|  | Identification / *Aitheantas* |  | Country / *Tír* | ISO country code / *Cód tíre ISO* |
|  | Commercial document reference / *Tagairt doiciméid tráchtála* |  |
| **I.18** | **Transport conditions / *Coinníollacha iompair*** | Ambient / *Comhthimpeallach* | Chilled / *Fuaraithe* | Frozen / *Reoite* |
| **I.19** | **Container number/Seal number / *Uimhir choimeádáin/Uimhir shéala*** |
|  | Container No / *Uimh. an choimeádáin* | Seal No / *Uimhir shéala* |  |
| **I.20** | **Certified as or for / *Deimhnithe mar nó le haghaidh*** |
|  | Further keeping / *Le coimeád a thuilleadh* |  |  |  |
|  |  | Confined establishment / *Bunaíocht theoranta* |  |  |
|  |  | Quarantine establishment / *Bunaíocht choraintín* |  |  |
|  |  |  | Other / *Eile* |  |
| **I.21** |  **For transit / *Le haghaidh iompair*** | **I.22** |  **For internal market / *Le haghaidh an mhargaidh inmheánaigh*** |
|  | Third country / *Tríú tír* | ISO country code / *Cód tíre ISO* | **I.23** |  |

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| **I.24** | **Total number of packages / *Líon iomlán pacáistí*** | **I.25** | **Total quantity / *Cainníocht*** ***iomlán*** | **I.26** | **Total net weight/gross weight (kg) / *Glanmheáchan/ollmheáchan (kg) iomlán*** |
| **I.27** | **Description of consignment / *Tuairisc ar an gcoinsíneacht*** |
| CN code / *Cód AC* | Species / *Speicis* | Subspecies/Category / *Fospeiceas/Catagóir* | Sex / *Inscne* | Identification system / *Córas aitheantais* | Identification number / *Uimhir aitheantais* | Age / *Aois* | Quantity / *Cainníocht* |
|  |  |  |  |  |  |  |  |
|  |  |  |  | Nature of commodity / *Cineál an tráchtearra* |  |  |  |
|  |  |  |  |  |  | Test / *Tástáil* |  |

| **COUNTRY / *TÍR***: Australia | **Certificate model CANIS-FELIS-FERRETS / *Deimhniú caighdeánach CANIS‑FELIS‑FERRETS*** |
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| **Part II: Certification / *Cuid II: Deimhniú*** | **II. Health information / *Faisnéis sláinte*** | **II.a**  | **Certificate reference / *Tagairt an deimhnithe*** | **II.b**  | **IMSOC reference / *Tagairt CBFRO*** |
| I, the undersigned official veterinarian hereby certify that the animals of the consignment described in Part I / *Deimhnímse, an tréidlia oifigiúil a bhfuil mo shíniú anseo thíos, leis seo an méid seo a leanas maidir le hainmhithe na coinsíneachta ar a dtugtar tuairisc i gCuid 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. tagann siad ó thríú tír nó ó chríoch, nó ó chrios de thríú tír nó de chríoch leis an gcód: AU - 0(1) atá, ar dháta eisiúna an deimhnithe sláinte ainmhithe seo, údaraithe maidir le teacht isteach san Aontas madraí, cat agus fíréad agus atá liostaithe i gCuid 1 d’Iarscríbhinn VIII a ghabhann le Rialachán Cur Chun Feidhme (AE) 2021/404 ón gCoimisiún;*(2) *either* [II.2. have been dispatched to the Union directly from the establishment of origin without passing through any other establishment;]*(2) cibé acu [II.2. seoladh ón mbunaíocht tionscnaimh go díreach chuig an Aontas iad gan dul trí aon bhunaíocht eile;]*(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) nó [II.2. rinneadh aon oibríocht tionóil amháin orthu sa tír thionscnaimh nó sa chríoch thionscnaimh, nó sa chrios den tír thionscnaimh nó den chríoch thionscnaimh sin, a tharla ar feadh tréimhse nach mó ná 6 lá i mbunaíocht lena gcomhlíontar na ceanglais seo a leanas:** *tá an bhunaíocht formheasta chun go ndéanfaidh an t‑údarás inniúil sa tríú tír nó sa chríoch oibríochtaí tionóil ar mhadraí, ar chait agus ar fhíréid i gcomhréir le hAirteagal 10 de Rialachán Tarmligthe (AE) 2019/2035 ón gCoimisiún;*
* *tá uimhir formheasa uathúil ag an mbunaíocht arna sannadh ag údarás inniúil an tríú tír nó na críche;*
* *tá an bhunaíocht liostaithe chun na críche sin ag údarás inniúil an tríú tír seolta nó na críche seolta chuig an Aontas, lena n‑áirítear an fhaisnéis a leagtar amach in Airteagal 21 de Rialachán Tarmligthe (AE) 2019/2035;*
* *comhlíonann an bhunaíocht na ceanglais maidir le taifid a choimeád dá bhforáiltear in Airteagal 73(2), pointe (a)(iv), de Rialachán Tarmligthe (AE) 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) nó [II.2. seoladh é/iad ó ionad dídine ainmhithe ina gcomhlíontar na ceanglais seo a leanas:* * *tá an bhunaíocht formheasta ag an údarás inniúil sa tríú tír nó sa chríoch i gcomhréir le hAirteagal 11 de Rialachán Tarmligthe (AE) 2019/2035;*
* *tá uimhir formheasa uathúil ag an mbunaíocht arna sannadh ag údarás inniúil an tríú tír nó na críche;*
* *tá sé liostaithe chun na críche sin ag údarás inniúil an tríú tír seolta nó na críche seolta, lena n‑áirítear an fhaisnéis dá bhforáiltear in Airteagal 21 de Rialachán Tarmligthe (AE) 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. luchtaíodh iad lena seoladh chuig an Aontas an \_\_\_/\_\_\_/\_\_\_\_ (ll/mm/bbbb) (4) i gcóir iompair a glanadh agus a díghalraíodh, roimh an luchtú, le díghalrán arna údarú ag údarás inniúil an tríú tír nó na críche agus arna tógáil ar bhealach a fhágann:** *nach féidir le hainmhithe éalú uaithi ná titim di;*
* *gur féidir iniúchadh amhairc a dhéanamh ar an spás ina gcoimeádtar na hainmhithe;*
* *go gcoisctear nó go n‑íoslaghdaítear éalú tuartha ainmhithe, bruscair nó beatha;]*

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 bhí siad faoi réir iniúchadh cliniciúil dá bhfuarthas toradh diúltach, arna dhéanamh ag tréidlia oifigiúil sa tríú tír nó sa chríoch thionscnaimh, nó sa chrios de thríú tír nó den chríoch thionscnaimh sin laistigh de na 48 uair an chloig dheireanacha roimh an am luchtaithe lena seoladh chuig an Aontas, chun comharthaí a bhrath lena léireofaí gur ann do ghalair, lena n‑áirítear na galair liostaithe dá dtagraítear in Iarscríbhinn I a ghabhann le Rialachán Tarmligthe (AE) 2020/692 is ábhartha don speiceas agus galair atá ag teacht chun cinn;*(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) cibé acu [II.5. beartaítear iad a thabhairt isteach go díreach sa Bhallstát cinn scríbe le cur ar leithlis:**(2) cibé acu [i mbunaíocht theoranta;]]**(2) nó [i mbunaíocht choraintín fhormheasta;]]*(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 )nó [II.5. a bhí 12 sheachtain d’aois ar a laghad ar dháta a vacsaínithe in aghaidh an chonfaidh agus go raibh 21 lá ar a laghad caite ón dáta a cuireadh an príomhvacsaíniú frithchonfaidh i gcrích (5) arna dhéanamh i gcomhréir leis na ceanglais bhailíochta a leagtar amach in Iarscríbhinn III a ghabhann le Rialachán (AE) Uimh. 576/2013 ó Pharlaimint na hEorpa agus ón gComhairle, agus go ndearnadh aon athvacsaíniú dá éis sin laistigh den tréimhse bhailíochta a bhain leis an vacsaíniú roimhe (6), agus:*(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) cibé acu [tagann siad ó thríú tír nó ó chríoch, agus i gcás idirthurais tá idirthuras sceidealaithe trí thríú tír nó trí chríoch a liostaítear in Iarscríbhinn II a ghabhann le Rialachán Cur Chun Feidhme (AE) Uimh. 577/2013 ón gCoimisiún agus soláthraítear mionsonraí an vacsaínithe/na vacsaínithe frithchonfaidh ábhartha i gcolúin 1 go 7 sa tábla thíos;]]*(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) nó [tagann siad ó thríú tír nó ó chríoch nó tá idirthuras sceidealaithe trí thríú tír nó trí chríoch nach liostaítear in Iarscríbhinn II a ghabhann le Rialachán Cur Chun Feidhme (AE) Uimh. 577/2013 ón gCoimisiún, agus:* (a) the details of the relevant anti-rabies vaccination(s) are provided in columns 1 to 7 in the table below / *soláthraítear mionsonraí an vacsaínithe/na vacsaínithe frithchonfaidh ábhartha i gcolún 1 go dtí colún 7 sa tábla thíos*, (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:]] / *tástáil le haghaidh toirtmheascadh antasubstainte (7), arna déanamh ar shampla fola arna thógáil ag an tréidlia arna údarú ag an údarás inniúil tráth nach lú ná 30 lá tar éis dháta an vacsaínithe roimhe agus 3 mhí ar a laghad roimh dháta eisiúna an deimhnithe sláinte ainmhithe sin, lenar cruthaíodh títear antasubstainte atá cothrom le nó níos mó ná 0,5 IU/ml (8) agus rinneadh aon athvacsaíniú dá éis sin laistigh den tréimhse bhailíochta a bhain leis an vacsaíniú roimhe, agus soláthraítear an dáta samplála chun an t‑imoibriú imdhíoneolaíoch a thástáil i gcolún 8 den tábla thíos:]]*

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| **Transponder / *Trasfhreagróir*** | **Date of vaccination [dd/mm/yyyy] / *Dáta vacsaínithe [ll/mm/bbbb]***  | **Name and manufacturer of vaccine / *Ainm agus monaróir na vacsaíne*** | **Batch number / *Baiscuimhir*** | **Validity of vaccination / *Bailíocht an vacsaínithe*** | **Date of blood sampling [dd/mm/yyyy] / *Dáta na samplála fola [ll/mm/bbbb]*** |
| **Alphanumeric code of the animal / *Cód alfa‑uimhriúil an ainmhí*** | **Date of implantation and/or reading (9) [dd/mm/yyyy] / *Dáta an inchlannaithe agus/nó an léimh (9) [ll/mm/bbbb]*** |
| **From [dd/mm/yyyy] / *Ó [ll/mm/bbbb]*** | **To [dd/mm/yyyy] / *Go dtí [ll/mm/bbbb]*** |
| 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) cibé acu [II.6. cuir madraí san áireamh a seolfar chuig Ballstát a liostaítear san Iarscríbhinn a ghabhann le Rialachán Cur Chun Feidhme (AE) 2018/878 ón gCoimisiún agus cóireáladh na madraí sin in aghaidh fíniú le Echinococcus multilocularis, agus soláthraítear mionsonraí na cóireála arna déanamh ag an tréidlia i gcomhréir le pointe 2 d’Iarscríbhinn XXI a ghabhann le Rialachán Tarmligthe (AE) 2020/692 (10) (11) sa tábla thíos:*

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| **Transponder or tattoo. Alphanumeric code of the dog / *Trasfhreagróir nó tatú. Cód alfa‑uimhriúil an mhadra*** | **Anti-Echinococcus treatment / *Cóireáil frith‑Echinococcus*** | **Administering veterinarian / *Tréidlia tabhartha*** |
| **Name and manufacturer of the product / *Ainm agus monaróir an táirge*** | **Date [dd/mm/yyyy] and time of treatment [00:00] / *Dáta [ll/mm/bbbb] agus am na cóireála [00:00]*** | **Name in capitals, stamp and signature / *Ainm i gceannlitreacha, stampa agus síniú*** |
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(2)*or* [II.6. include dogs which have not been treated against infestation with *Echinococcus multilocularis.*]*(2) nó [II.6. lena n‑áirítear madraí nár cóireáladh in aghaidh a bhfínithe le Echinococcus multilocularis.]*(2)*or* [II.6. include dogs destined for direct entry into the Member State of destination to be isolated in:*(2) nó [II.6. áirigh madraí a bheartaítear teacht isteach go díreach sa Bhallstát cinn scríbe le cur ar leithlis:*(1) *either* [a confined establishment.]]*(1) cibé acu [i mbunaíocht theoranta.]]*(1) *or* [an approved quarantine establishment.]]*(1) nó [i mbunaíocht choraintín fhormheasta.]]***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.***Nótaí:****Tá an deimhniú sláinte ainmhithe seo beartaithe le haghaidh theacht isteach san Aontas madraí, cat agus fíréad ar bhonn tráchtála, lena n‑áirítear i gcásanna ina bhfuil siad le dul chuig bunaíocht theoranta nó chuig bunaíocht choraintín fhormheasta agus i gcásanna nach é an tAontas ceann scríbe críochnaitheach na n‑ainmhithe agus maidir le teacht isteach san Aontas madraí, cat agus fíréad arna n‑aistriú i gcomhréir le hAirteagal 5(4) de Rialachán (AE) Uimh. 576/2013 ó Pharlaimint na hEorpa agus ón gComhairle.**I gcomhréir leis an gComhaontú maidir le Ríocht Aontaithe na Breataine Móire agus Thuaisceart Éireann a bheith ag tarraingt siar as an Aontas Eorpach agus as an gComhphobal Eorpach do Fhuinneamh Adamhach, agus go háirithe Airteagal 5(4) den Phrótacal maidir le hÉirinn/Tuaisceart Éireann i dteannta le hIarscríbhinn 2 a ghabhann leis an bPrótacal sin, áirítear an Ríocht Aontaithe i ndáil le Tuaisceart Éireann sna tagairtí don Aontas sa deimhniú sláinte ainmhithe seo.**Déanfar an deimhniú sláinte ainmhithe seo a chomhlánú i gcomhréir leis na nótaí a bhaineann le comhlíonadh na ndeimhnithe dá bhforáiltear i gCaibidil 4 d’Iarscríbhinn I a ghabhann le Rialachán Cur Chun Feidhme (AE) 2020/2235 ón gCoimisiún.***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.

***Cuid I:****Tagairt bhosca I.20: Deimhnithe mar nó le haghaidh: Sonraigh:* * *‘Le coimeád a thuilleadh’ i gcás ina n‑aistrítear madraí, cait nó fíréid i gcomhréir le Teideal V de Chuid II de Rialachán Tarmligthe (AE) 2020/692;*
* *Bunaíocht theoranta: mar a shainmhínítear in Airteagal 4(48) de Rialachán (AE) 2016/429 ó Pharlaimint na hEorpa agus ón gComhairle;*
* *Bunaíocht choraintín fhormheasta: mar a shainmhínítear in Airteagal 3(9) de Rialachán Tarmligthe (AE) 2020/688 ón gCoimisiún;*
* *‘ainmhithe eile’ i gcás ina n‑aistrítear madraí (Canis lupus familiaris), cait (Felis silvestris catus) nó firéid (Mustela putorius furo) i gcomhréir le hAirteagal 5(4) de Rialachán (AE) Uimh. 576/2013 ó Pharlaimint na hEorpa agus ón gComhairle.*

**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.***Cuid II:****(1) Cód an chreasa mar atá sé le feiceáil i gcolún 2 den tábla i gCuid 1 d’Iarscríbhinn VIII a ghabhann le Rialachán Cur Chun Feidhme (AE) 2021/404.**(2) Scrios murab infheidhme.**(3) Níl sé infheidhme maidir le gluaiseacht madraí, cat agus firéad seachas gluaiseachtaí neamhthráchtálacha, a choimeádtar mar pheataí i dteaghlaigh nach féidir a dhéanamh i gcomhréir leis na coinníollacha a leagtar síos in Airteagal 245(2) nó in Airteagail 246(1) agus (2) de Rialachán (AE) 2016/429.**(4) Dáta luchtaithe: ní fhéadfaidh sé a bheith ar dháta roimh dháta údaraithe an chreasa maidir le teacht isteach san Aontas, ná i dtréimhse a raibh bearta sriantacha i bhfeidhm ag an Aontas lena linn in aghaidh theacht isteach na n‑ainmhithe sin ón gcrios sin.**(5) Measfar aon athvacsaíniú a bheith ina vacsaíniú príomhúil mura ndearnadh é laistigh den tréimhse bhailíochta a bhain le vacsaíniú roimhe.**(6) Cuirfear cóip dhílis dheimhnithe de mhionsonraí aitheantais agus vacsaínithe na n‑ainmhithe lena mbaineann i gceangal leis an deimhniú sláinte ainmhithe.**(7) An tástáil le haghaidh thoirtmheascadh antasubstainte an chonfaidh dá dtagraítear i bpointe II.5.:**- déanfar í ar shampla arna bhailiú ag tréidlia arna údarú ag an údarás inniúil, 30 lá ar a laghad tar éis dháta an vacsaínithe agus 3 mhí roimh dháta an tseolta chuig an Aontas;**- tomhaisfidh sé leibhéal antasubstainte neodraithe ar víreas an chonfaidh i séiream a bheidh comhionann le nó níos mó ná 0,5 IU/ml;**- déanfaidh saotharlann oifigiúil í;**- ní dhéanfar é a athnuachan maidir le hainmhí, tar éis na tástála sin dá bhfuarthas torthaí sásúla, a athvacsaíníodh in aghaidh an chonfaidh laistigh den tréimhse bhailíochta a bhain le vacsaíniú roimhe.* *Cuirfear cóip dhílis dheimhnithe den tuarascáil oifigiúil ón tsaotharlann oifigiúil maidir le toradh na tástála antasubstainte confaidh dá dtagraítear i bpointe II.5. i gceangal leis an deimhniú sláinte ainmhithe.**(8) Tríd an toradh sin a dheimhniú, dearbhaíonn an tréidlia oifigiúil gur fhíoraigh sé, ar feadh a chumais agus i gcás inar gá le teagmhálaithe leis an tsaotharlann a shonraítear sa tuarascáil, barántúlacht thuarascáil na saotharlainne maidir le torthaí na tástála maidir le toirtmheascadh antasubstainte dá dtagraítear i bpointe II.5.**(9) I gcomhar le nóta (6), déanfar marcáil na n‑ainmhithe lena mbaineann trí thrasfhreagróir a ionchlannú a fhíorú sula ndéanfar aon iontráil sa deimhniú sláinte ainmhithe sin agus déanfar an mharcáil sin roimh aon vacsaíniú i gcónaí, nó i gcás inarb infheidhme, roimh aon tástáil dá ndéantar ar na hainmhithe sin.**(10) Maidir leis an gcóireáil in aghaidh fíniú le Echinococcus multilocularis dá dtagraítear i bpointe II.6., ní mór:**- í a bheith á tabhairt ag tréidlia laistigh de thréimhse nach faide ná 48 n‑uair an chloig agus a chríochnaíonn tráth nach lú ná 24 uair an chloig roimh an am sceidealaithe chun na madraí a sheoladh chuig ceann de na Ballstáit nó chuig codanna díobh a liostaítear san Iarscríbhinn a ghabhann le Rialachán Cur Chun Feidhme (AE) 2018/878 ón gCoimisiún;**- atá comhdhéanta de tháirge íocshláinte formheasta ina bhfuil an dáileog iomchuí de praziquantel nó de shubstaint atá gníomhach ó thaobh na cógaseolaíochta de, lenar cruthaíodh, ina n‑aonar nó i gcomhar le chéile, go laghdaítear leo ualaí na bhfoirmeacha stéigeacha aibí agus neamhaibí de Echinococcus multilocularis sa speiceas óstála lena mbaineann.**(11) Déanfar an tábla dá dtagraítear i bpointe II.6 a úsáid chun mionsonraí cóireála breise a dhoicimeadú má thugtar an chóireáil tar éis an dáta a síníodh an deimhniú sláinte ainmhithe agus roimh theacht isteach sceidealaithe i gceann de na Ballstáit nó i gcodanna díobh a liostaítear san Iarscríbhinn a ghabhann le Rialachán Cur Chun Feidhme (AE) 2018/878.* |
| **Official veterinarian / *Tréidlia oifigiúil*** |
| Name (in capital letters) / *Ainm (i gceannlitreacha)* |  |  |  |
| Date / *Dáta* |  | Qualification and title / *Cáilíocht agus teideal* |  |
| Stamp / *Stampa* |  | Signature / *Síniú* |  |