

Animal health certificate for the non-commercial movement into a Member State from a territory or third country of dogs, cats or ferrets
in accordance with Article 5(1) and (2) of Regulation (EU) No 576/2013 /
Veterinarskega spričevala za netrgovske premike psov, mačk ali belih dihurjev v državo
članico z ozemlja ali iz tretje države
v skladu s členom 5(1) in (2) Uredbe (EU) št. 576/2013

United Kingdom /
Velika Britanija

Veterinary certificate to EU /
Veterinarsko spričevalo Evropski uniji

Part I : Details of dispatched consignment Del I: Podrobnosti odpromljenje pošiljke	I.1. Consignor / Pošiljatelj Name / Ime Address / Naslov Tel. / Telefon				I.2. Certificate reference No / Referenčna številka spričevala I.3. Central competent authority / Osrednji pristojni organ DEFRA		I.2.a I.4. Local competent authority / Lokalni pristojni organ	
	I.5. Consignee / Prejemnik Name / Ime Address / Naslov Postal code / Poštna številka Tel. / Telefon				I.6. Person responsible for the consignment in the EU / Oseba v EU, odgovorna za pošiljko			
	I.7. Country of origin / Država porekla United Kingdom	ISO Code / Oznaka ISO GB	I.8. Region of origin / Regija izvora	Code Oznaka	I.9. Country of destination / Namembna država	ISO Code / Oznaka ISO	I.10 Region of destination / Namembna regija	Code/ Oznaka
	I.11. Place of origin / Kraj izvora				I.12. Place of destination / Namembni kraj			
	I.13. Place of loading / Kraj natovarjanja				I.14. Date of departure / Datum pošiljanja			
	I.15. Means of transport / Prevozno sredstvo				I.16. Entry BIP in EU / Mejna kontrolna točka vstopa v EU			
	I.18. Description of commodity / Opis blaga				I.19. Commodity code (HS code) / Oznaka blaga (oznaka HS) 010619			
	I.21. Temperature of products / Temperatura proizvodov				I.22. Total number of packages / Skupno število pakiranj			
	I.23. Seal/Container No / Številka zalivke/kontejnerja				I.24. Type of packaging / Vrsta pakiranja			
	I.25. Commodities certified for: / Blago s spričevalom za: Pets / Hišne živali <input type="checkbox"/>							
	I.26. For transit to 3 rd Country / Za tranzit v tretjo državo				I.27. For import or admission into EU / Za uvoz ali vstop v EU			
	I.28. Identification of the commodities / Identifikacija blaga Species (Scientific name) / Sex / Colour / Breed / Identification number / Vrsta (znanstveno ime) Spol Barva Pasma Identifikacijska številka				Identification system / Identifikacijski sistem Date of birth / [dd/mm/yyyy] Datum rojstva [dd/mm/llll]			

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II.	Health information / Potrdilo o zdravstvenem stanju	II.a. Certificate reference No / Referenčna številka spričevala	II.b.
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I, the undersigned official veterinarian⁽¹⁾/veterinarian authorised by the competent authority⁽¹⁾ of the United Kingdom, / Spodaj podpisani uradni veterinar⁽¹⁾/veterinari, pooblaščen s strani pristojnega organa⁽¹⁾ Velika Britanija, certify that / potrjujem, da:

Purpose/nature of journey attested by the owner / Namen/Narava potovanja, ki ga/jo potrdi lastnik:

II.1. the attached declaration⁽²⁾ by the owner or the natural person who has authorisation in writing from the owner to carry out the non-commercial movement of the animals on behalf of the owner, supported by evidence⁽³⁾, states that the animals described in Box I.28 will accompany the owner or the natural person who has authorisation in writing from the owner to carry out the non-commercial movement of the animals on behalf of the owner within not more than five days of his movement and are not subject to a movement that aims at their sale or a transfer of ownership, and during the non-commercial movement will remain under the responsibility of / v priloženi izjavi⁽²⁾ lastnika ali fizične osebe, ki ima pisno dovoljenje lastnika za opravljanje netrgovskega premika živali v imenu lastnika, podprt z dokazi⁽³⁾, je navedeno, da bodo živali, opisane v rubriki I.28, spremljale lastnika ali fizično osebo, ki ima pisno dovoljenje lastnika, da opravlja netgovski premik živali v imenu lastnika, najpozneje v petih dneh od njegovega premika in da cilj premika ni njihova prodaja ali prenos lastništva ter da bo med netrgovskim premikom zanje še naprej odgovoren

⁽¹⁾either/bodisi [the owner;] / [lastnik;]

⁽¹⁾or/bodisi [the natural person who has authorisation in writing from the owner to carry out the non-commercial movement of the animals on behalf of the owner;] / [fizična oseba, ki ima pisno dovoljenje lastnika, da opravi netgovski premik živali v imenu lastnika;]

⁽¹⁾or/ bodisi [the natural person designated by a carrier contracted by the owner to carry out the non-commercial movement of the animals on behalf of the owner;] / [fizična oseba, ki jo določi prevoznik, s katerim je lastnik sklenil pogodbo za opravljanje netrgovskega premika živali v imenu lastnika;]

⁽¹⁾ either/bodisi [II.2. the animals described in Box I.28 are moved in a number of five or less;] / [gre za premik pet ali manj živali iz rubrike I.28;]

⁽¹⁾ or/ bodisi [II.2. the animals described in Box I.28 are moved in a number of more than five, are more than six months old and are going to participate in competitions, exhibitions or sporting events or in training for those events, and the owner or the natural person referred to in point II.1 has provided evidence⁽³⁾ that the animals are registered / gre za premik več kot pet živali iz rubrike I.28, ki so starejše od šest mesecev in se bodo udeležile tekmovanj, razstav ali športnih prireditev ali usposabljanja za navedene dogodke, lastnik ali fizična oseba iz točke II.1 pa je predložil oz. predložila dokaze⁽³⁾, da so živali registrirane

⁽¹⁾ either/bodisi [to attend such an event;] / [za udeležbo na takšnem dogodku;]

⁽¹⁾ or/ bodisi [with an association organising such events;] / [pri združenju, ki takšne dogodke organizira;]

Attestation of rabies vaccination and rabies antibody titration test / Potrdilo o cepljenju proti steklini in test titracije protiteles proti steklini:

⁽¹⁾ either/bodisi [II.3. the animals described in Box I.28 are less than 12 weeks old and have not received an anti-rabies vaccination, or are between 12 and 16 weeks old and have received an anti-rabies vaccination, but 21 days at least have not elapsed since the completion of the primary vaccination against rabies carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013⁽⁴⁾, and / živali iz rubrike I.28 so mlajše od 12 tednov in niso cepljene proti steklini ali so stare med 12 in 16 tednov in so bile cepljene proti steklini, vendar ni minilo vsaj 21 dni od zaključka primarnega cepljenja proti steklini, ki se opravlja v skladu z zahtevami o veljavnosti iz Priloge III k Uredbi (EU) št. 576/2013⁽⁴⁾, ter

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	II.3.1 the territory or third country of provenance of the animals indicated in Box I.1 is listed in Annex II to Implementing Regulation (EU) No 577/2013 and the Member State of destination indicated in Box I.5 has informed the public that it authorises the movement of such animals into its territory, and they are accompanied by / je provenienčno ozemlje ali provenienčna tretja država živali iz rubrike I.1 na seznamu v Prilogi II k Izvedbeni uredbi (EU) št. 577/2013, namembna država članica iz rubrike I.5 pa je obvestila javnost, da dovoljuje premik takšnih živali na njeno ozemlje, in jih spremlja ⁽¹⁾ either/bodisi [II.3.2 the attached declaration(5) of the owner or the natural person referred to in point II.1 stating that from birth until the time of the non-commercial movement the animals have had no contact with wild animals of species susceptible to rabies;] / priložena izjava ⁽⁵⁾ lastnika ali fizične osebe iz točke II.1, v kateri je navedeno, da živali od rojstva do trenutka netrgovskega premika niso bile v stiku z divjimi živalmi vrst, ki so dovzetne za steklino;] ⁽¹⁾ either/bodisi [II.3.2 their mother, on whom they still depend, and it can be established that the mother received before their birth an anti-rabies vaccination which complied with the validity requirements set out in Annex III to Regulation (EU) No 576/2013;] / njihova mati, od katere so še vedno odvisne, in je mogoče ugotoviti, da je mati bila cepljena proti steklini pred njihovim rojstvom s cepivom, ki izpolnjuje zahteve glede veljavnosti iz Priloge III k Uredbi (EU) št. 576/2013;] ⁽¹⁾ and/or/ bodisi/in [II.3.2 the animals described in Box I.28 were at least 12 weeks old at the time of vaccination against rabies and at least 21 days have elapsed since the completion of the primary anti-rabies vaccination ⁽⁴⁾ carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 and any subsequent revaccination was carried out within the period of validity of the preceding vaccination ⁽⁶⁾ ; and / živali iz rubrike I.28 so stare vsaj 12 tednov v trenutku cepljenja proti steklini in je minilo najmanj 21 dni od zaključka primarnega cepljenja proti steklini ⁽⁴⁾ , opravljenega v skladu z zahtevami o veljavnosti iz Priloge III k Uredbi (EU) št. 576/2013, vsa nadaljnja ponovna cepljenja pa so bila opravljena v obdobju veljavnosti predhodnega cepljenja ⁽⁶⁾ ; ter ⁽¹⁾ either/bodisi [II.3.1 the animals described in Box I.28 come from a territory or a third country listed in Annex II to Implementing Regulation (EU) No 577/2013, either directly, through a territory or a third country listed in Annex II to Implementing Regulation (EU) No 577/2013 or through a territory or a third country other than those listed in Annex II to Implementing Regulation (EU) No 577/2013 in accordance with point (c) of Article 12(1) of Regulation (EU) No 576/2013 ⁽⁷⁾ , and the details of the current anti-rabies vaccination are provided in the table below;] / živali iz rubrike I.28 prihajajo z ozemlja ali iz tretje države s seznama v Prilogi II k Izvedbeni uredbi (EU) št. 577/2013 bodisi neposredno ali preko ozemlja ali tretje države s seznama v Prilogi II k Izvedbeni uredbi (EU) št. 577/2013 ali prek ozemlja ali tretje države, ki ni na seznamu v Prilogi II k Izvedbeni uredbi (EU) št. 577/2013, v skladu s točko (c) člena 12(1) Uredbe (EU) št. 576/2013 ⁽⁷⁾ , podrobnosti o trenutnem cepljenju proti steklini pa so navedene v spodnji tabeli;] ⁽¹⁾ or/bodisi [II.3.1 the animals described in Box I.28 come from, or are scheduled to transit through, a territory or third country other than those listed in Annex II to Implementing Regulation (EU) No 577/2013 and a rabies antibody titration test ⁽⁸⁾ , carried out on a blood sample taken by the veterinarian authorised by the competent authority on the date indicated in the table below not less than 30 days after the preceding vaccination and at least three months prior to the date of issue of this certificate, proved an antibody titre equal to or greater than 0.5 IU/ml ⁽⁹⁾ and any subsequent revaccination was carried out within the period of validity of the preceding vaccination ⁽⁶⁾ , and the details of the current anti-rabies vaccination and the date of sampling for testing the immune response are provided in the table below: / živali iz rubrike I.28 prihajajo z ozemlja ali iz tretje države ali so načrtovane za tranzit preko ozemlja ali tretje države, ki ni na seznamu v Prilogi II k Izvedbeni uredbi (EU) št. 577/2013, s testom titracije protiteles proti steklini ⁽⁸⁾ , opravljenem na vzorcu krvi, ki ga je veterinar, pooblaščen s strani pristojnega organa, odvzel na datum, naveden v spodnji tabeli, ne manj kot 30 dni po predhodnem cepljenju in najmanj tri mesece pred datumom izdaje tega spričevala, je bil dokazan titer protiteles višini vsaj 0,5 IE/ml ⁽⁹⁾ , vsakršno pozneje ponovno cepljenje je bilo opravljeno v obdobju veljavnosti predhodnega cepljenja ⁽⁶⁾ , podrobnosti o trenutnem cepljenju proti steklini in datum vzorčenja za testiranje imunskega odziva pa so navedeni v spodnji tabeli:]		

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Transponder or tattoo / Transponderja ali vtetovirano znamenje		Date of implantation and/or reading ⁽¹⁰⁾ [dd/mm/yyyy] Datum vsaditve in/ali odčitanja ⁽¹⁰⁾ [dd/mm/llll]	Name and manufacturer of vaccine / Ime in proizvajalec cepiva	Batch number / Serijska številka	Validity of vaccination / Veljavnost cepljenja	Date of the blood sampling [dd/mm/yyyy] / Datum vzorčenja krvi [dd/mm/llll]
					From [dd/mm/yyyy] / od [dd/mm/llll]	

Attestation of anti-parasite treatment / Potrjevanje zdravljenja zaradi parazitov:

- ⁽¹⁾either/bodisi [II.4. the dogs described in Box I.28 are destined for a Member State listed in the Annex to Commission Implementing Regulation (EU) 2018/878 and have been treated against Echinococcus multilocularis, and the details of the treatment carried out by the administering veterinarian in accordance with Article 6 of Commission Delegated Regulation (EU) 2018/772 ⁽¹¹⁾⁽¹²⁾⁽¹³⁾ are provided in the table below. / psi iz rubrike I.28 so namenjeni v državo članico s seznama v Prilogi k izvedbeni uredbi Komisije (EU) 2018/878 in so bili zdravljeni zaradi Echinococcus multilocularis, podrobnosti o zdravljenju, ki ga je opravil veterinar v skladu s členom 6 delegirane uredbe Komisije (EU) 2018/772 ⁽¹¹⁾⁽¹²⁾⁽¹³⁾, pa so navedeni v spodnji tabeli.]
- ⁽¹⁾or/bodisi [II.4. the dogs described in Box I.28 have not been treated against *Echinococcus multilocularis*⁽¹¹⁾. / psi iz rubrike I.28 niso bili zdravljeni zaradi *Echinococcus multilocularis*⁽¹¹⁾.]

Transponder or tattoo number of the dog / Številka transponderja ali vtetoviranega znamenja psa	Anti-echinococcus treatment / Zdravljenje zaradi echinokoka		Administering veterinarian / Lečeči veterinar
	Name and manufacturer of the product / Ime in proizvajalec zdravila	Date [dd/mm/yyyy] and time of treatment [00:00] / Datum [dd/mm/llll] in čas zdravljenja [00:00]	

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Notes / Opombe

- a) This certificate is meant for dogs (*Canis lupus familiaris*), cats (*Felis silvestris catus*) and ferrets (*Mustela putorius furo*). / To spričevalo se uporablja za pse (*Canis lupus familiaris*), mačke (*Felis silvestris catus*) in bele dihurje (*Mustela putorius furo*).
 - b) This certificate is valid for 10 days from the date of issue by the official veterinarian until the date of the documentary and identity checks at the designated Union travellers' point of entry, available at http://ec.europa.eu/food/animal/liveanimals/pets/pointsentry_en.htm / To spričevalo velja 10 dni od datuma izdaje s strani uradnega veterinarja do datuma pregledov dokumentov in identitete na določeni vstopni točki potnikov v Unijo (na voljo na http://ec.europa.eu/food/animal/liveanimals/pets/pointsentry_en.htm).
- In the case of transport by sea, that period of 10 days is extended by an additional period corresponding to the duration of the journey by sea. / V primeru prevoza po morju se navedeno obdobje 10 dni podaljša za dodatno obdobje, ki ustreza trajanju potovanja po morju.
- For the purpose of further movement into other Member States, this certificate is valid from the date of the documentary and identity checks for a total of four months or until the date of expiry of the validity of the anti-rabies vaccination or until the conditions relating to animals less than 16 weeks old referred to in point II.3 cease to apply, whichever date is earlier. Please note that certain Member States have informed that the movement into their territory of animals less than 16 weeks old referred to in point II.3 is not authorised. You may wish to inquire at http://ec.europa.eu/food/animal/liveanimals/pets/index_en.htm / Za nadaljnje premike v druge države članice to spričevalo velja od dneva pregledov dokumentacije in identitete za skupaj štiri mesece ali do izteka veljavnosti cepljenja proti steklini ali dokler se pogoj, ki se nanašajo na živali iz točke II.3, mlajše od 16 tednov, prenehajo uporabljati, pri čemer velja zgodnejši datum. Upoštevajte, da so nekatere države članice sporočile, da premik živali iz točke II.3, mlajših od 16 tednov, na njihovo ozemlje ni dovoljen. Več informacij na http://ec.europa.eu/food/animal/liveanimals/pets/index_en.htm.

Part I / Del I :

Box 1.5: / Rubrika I.5.: *Consignee*: indicate Member State of first destination. / *Prejemnik*: navedite državo članico prvega namembnega kraja.

Box 1.28: / Rubrika I.28: *Identification system*: select of the following: transponder or tattoo. *Identification number*: indicate the transponder or tattoo alphanumeric code. *Date of birth/breed*: as stated by the owner. / *Identifikacijski sistem*: izberite: transponder ali vtetovirano znamenje. *Identifikacijska številka*: navedite črkovno-številčno oznako transponderja ali vtetoviranega znamenja. *Datum rojstva/pasma*: po navedbi lastnika.

Part II / Del II:

- (1) Keep as appropriate / Neustrezno črtati.
- (2) The declaration referred to in point II.1 shall be attached to the certificate and comply with the model and additional requirements set out in Part 3 of Annex IV to Implementing Regulation (EU) No 577/2013. / Izjava iz točke II.1 se priloži spričevalu, izpolnjuje pa zahteve za vzorec in dodatne zahteve iz dela 3 Priloge IV k Izvedbeni uredbi (EU) št. 577/2013.
- (3) The evidence referred to in point II.1 (e.g. boarding pass, flight ticket) and in point II. 2 (e.g. receipt of entry to the event, proof of membership) shall be surrendered on request by the competent authorities responsible for the checks referred to in point (b) of the Notes. / Dokazi iz točke II.1 (npr. vstopni kupon, letalska vozovnica) in točke II.2 (npr. potrdilo o udeležbi na prireditvi, dokaz o članstvu) se na zahtevo predajo pristojnim organom, odgovornim za preglede iz točke (b) Opomb.
- (4) Any revaccination must be considered a primary vaccination if it was not carried out within the period of validity of a previous vaccination. / Kakršno koli obnovitveno cepljenje se šteje za primarno cepljenje, če ni opravljeno v obdobju veljavnosti predhodnega cepljenja.
- (5) The declaration referred to in point II.3.2 to be attached to the certificate complies with the format, layout and language requirements laid down in Parts 1 and 3 of Annex I to Implementing Regulation (EU) No 577/2013. / Izjava iz točke II.3.2 se priloži spričevalu, izpolnjuje pa zahteve o formatu, obliku in jeziku iz delov 1 in 3 Priloge I k Izvedbeni uredbi (EU) št. 577/2013.

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(6)	A certified copy of the identification and vaccination details of the animals concerned shall be attached to the certificate. / Spričevalu se priloži overjena kopija z identifikacijo in podrobnostmi cepljenja zadevnih živali.		
(7)	The third option is subject to the condition that the owner or the natural person referred to in point II.1 provides, on request by the competent authorities responsible for the checks referred to in point (b), a declaration stating that the animals have had no contact with animals of species susceptible of rabies and remain secure within the means of transport or the perimeter of an international airport during the transit through a territory or a third country other than those listed in Annex II to Implementing Regulation (EU) No 577/2013. This declaration shall comply with the format, layout and language requirements set out in Parts 2 and 3 of Annex I to Implementing Regulation (EU) No 577/2013. / Za tretjo možnost velja pogoj, da lastnik ali fizična oseba iz točke II.1 na zahtevo pristojnih organov, odgovornih za preglede iz točke (b), predloži izjavo, v kateri navede, da živali niso bile v stiku z živalskimi vrstami, ki so dovezene za steklino, in so bile zavarovane v prevoznem sredstvu ali znotraj območja mednarodnega letališča med tranzitom preko ozemlja ali tretje države, ki ni na seznamu v Prilogi II k Izvedbeni uredbi (EU) št. 577/2013. Ta izjava je skladna s formatom, obliko in jezikovnimi zahtevami iz delov 2 in 3 Priloge I k Izvedbeni uredbi (EU) št. 577/2013.		
(8)	<p>The rabies antibody titration test referred to in point II.3.1 / Test titracije protiteles proti steklini iz točke II.3.1:</p> <ul style="list-style-type: none"> - must 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 three months before the date of import; / opravljen mora biti na vzorcu, ki ga odvzame veterinar, pooblaščen s strani pristojnega organa, vsaj 30 dni po datumu cepljenja in tri mesece pred datumom uvoza; - must measure a level of neutralising antibody to rabies virus in serum equal to or greater than 0.5 IU/ml; / mora izmeriti stopnjo nevtralizacijskih protiteles proti steklini v serumu, ki mora znašati vsaj 0.5 IE/ml; - must be performed by a laboratory approved in accordance with Article 3 of Council Decision 2000/258/EC (list of approved laboratories available at: http://ec.europa.eu/food/animal/liveanimals/pets/approval_en.htm) / mora opraviti laboratorij, odobren v skladu s členom 3 Odločbe Sveta 2000/258/ES (seznam odobrenih laboratorijskih je na voljo na http://ec.europa.eu/food/animal/liveanimals/pets/approval_en.htm); - does not have to 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. / ni treba ponoviti pri živali, ki je bila po testu z zadovoljivimi rezultati ponovno cepljena proti steklini v obdobju veljavnosti predhodnega cepljenja. <p>A certified copy of the official report from the approved laboratory on the results of the rabies antibody test referred to in point II.3.1 shall be attached to the certificate / Overjena kopija uradnega poročila iz odobrenega laboratorijskega rezultatov testa na protitelesa proti steklini iz točke II.3.1 se priloži spričevalu.</p>		
(9)	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.3.1 / Z overitvijo teh rezultatov uradni veterinar potrjuje, da je na najboljši možen način in po potrebi v stiku z laboratorijskim, navedenim v poročilu, preveril avtentičnost laboratorijskega poročila o rezultatih testa titracije protiteles iz točke II.3.1.		
(10)	In conjunction with footnote (6), the marking of the animals concerned by the implantation of a transponder or by a clearly readable tattoo applied before 3 July 2011 must be verified before any entry is made in this certificate and must always precede any vaccination, or where applicable, testing carried out on those animals / V povezavi z opombo (6) je treba označevanja zadevnih živali z vsaditvijo transponderja ali z jasno čitljivim znamenjem, ki so bila izvedena pred 3. julijem 2011, preveriti pred vsakim vnosom v to spričevalo in jih je treba vedno opraviti pred vsakim cepljenjem, ali, kadar je to primerno, testiranjem, opravljenim na navedenih živalih.		
(11)	The treatment against <i>Echinococcus multilocularis</i> referred to in point II.4 must / Zdravljenje zaradi <i>Echinococcus multilocularis</i> iz točke II.4 mora:		
	<ul style="list-style-type: none"> - be administered by a veterinarian within a period of not more than 120 hours and not less than 24 hours before the time of the scheduled entry of the dogs into one of the Member States or parts thereof listed in the Annex to Implementing Regulation (EU) 2018/878 / opraviti veterinar v obdobju največ 120 ur in najmanj 24 ur pred načrtovanim vstopom psov v eno od držav članic ali njihovih delov s seznama v Prilogi k izvedbeni uredbi (EU) 2018/878; 		

Animal health certificate for the non-commercial movement into a Member State from a territory or third country of dogs, cats or ferrets

in accordance with Article 5(1) and (2) of Regulation (EU) No 576/2013 /

**Veterinarskega spričevala za netrgovske premike psov, mačk ali belih dihurjev v državo
članico z ozemlja ali iz tretje države
v skladu s členom 5(1) in (2) Uredbe (EU) št. 576/2013**

**United Kingdom /
Velika Britanija**

**Veterinary certificate to EU /
Veterinarsko spričevalo Evropski uniji**

II.	Health information/ Potrdilo o zdravstvenem stanju	II.a. Certificate reference No / Referenčna številka spričevala	II.b.
- 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 <i>Echinococcus multilocularis</i> in the host species concerned / pri njem se mora uporabiti odobreno zdravilo, ki vsebuje ustrezno dozo prazikvantela ali farmakološko aktivnih snovi, ki same ali skupaj dokazano zmanjšujejo obremenitev z odraslimi in nezreliimi črevesnimi oblikami <i>Echinococcus multilocularis</i> pri zadevnih gostiteljskih vrstah.			
(12)		The table referred to in point II.4 must be used to document the details of a further treatment if administered after the date the 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 / Tabelo iz točke II.4 je treba uporabiti za dokumentiranje podrobnosti o nadaljnjem zdravljenju, če se opravi po datumu podpisa spričevala in pred načrtovanim vstopom v eno od držav članic ali njihovih delov s seznama v Prilogi k izvedbeni uredbi (EU) 2018/878.	
(13)		The table referred to in point II.4 must be used to document the details of treatments if administered after the date the certificate was signed for the purpose of further movement into the United Kingdom or Member States described in point (b) of the Notes and in conjunction with footnote (11) / Tabelo iz točke II.4 je treba uporabiti za dokumentiranje podrobnosti o zdravljenju, če se opravi po datumu podpisa spričevala za nadaljnje premike v druge države članice iz točke (b) v opombah in v povezavi z opombo (11).	

**Animal health certificate for the non-commercial movement into a Member State from a territory or third country of dogs, cats or ferrets
in accordance with Article 5(1) and (2) of Regulation (EU) No 576/2013 /
Veterinarskega spričevala za netrgovske premike psov, mačk ali belih dihurjev v državo
članico z ozemlja ali iz tretje države
v skladu s členom 5(1) in (2) Uredbe (EU) št. 576/2013**

**United Kingdom /
Velika Britanija**

**Veterinary certificate to EU /
Veterinarsko spričevalo Evropski uniji**

II. Health information/ Potrdilo o zdravstvenem stanju	II.a. Certificate reference No / Referenčna številka spričevala	II.b.
Official veterinarian/Authorised veterinarian / Uradni veterinar/pouplaščeni veterinar		
Name (in capital letters) / Ime (s tiskanimi črkami):	Qualification and title / Izobrazba in naziv:	
Address / Naslov:		
Telephone / Telefon:		
Date / Datum:	Signature / Podpis:	
Stamp / Žig:		
Endorsement by the competent authority (not necessary when the certificate is signed by an official veterinarian) / Potrditev pristojnega organa (ni potrebna, kadar spričevalo podpiše uradni veterinar)		
Name (in capital letters) / Ime (s tiskanimi črkami):	Qualification and title / Izobrazba in naziv:	
Address / Naslov:		
Telephone / Telefon:		
Date / Datum:	Signature / Podpis:	
Stamp / Žig:		
Official at the travellers' point of entry (for the purpose of further movement into other Member States) / Uradnik na vstopni točki potnikov (za nadaljnje premike v druge države članice)		
Name (in capital letters) / Ime (s tiskanimi črkami):	Title / Naziv:	
Address / Naslov:		
Telephone / Telefon:		
E-mail address / E-naslov:		
Date of completion of documentary and identity checks / Datum zaključka pregledov dokumentacije in istovetnosti:		
Signature / Podpis:		Stamp / Žig:

This document is valid for re-entry to the United Kingdom from EU Member States for four months after the date of issue by the Official Veterinarian/(Section II).

Where required, the *Echinococcus multilocularis* treatment should be recorded in section II.4 of the certificate before return to the United Kingdom. This treatment must be administered not less than 24 hours and not more than 120 hours before the pet is landed in the United Kingdom. A practice stamp may be used in place of an Official Veterinarian's/authorised veterinarian's stamp in II.4.

A record of any further rabies vaccination carried out while in the EU should accompany this certificate.

Please note if a young pet is authorised for travel out of Great Britain in accordance with Article 11 of Regulation 576/2013, it will not be able to return until it has received an anti-rabies vaccination that complies with the validity requirements of Annex III of Regulation 576/2013. In practice this means your pet must be at least 15 weeks old to enter Great Britain.

Written declaration referred to in Article 25(3) of Regulation (EU) No 576/2013 / **Pisna izjava** iz člena 25(3) Uredbe (EU) št. 576/2013

Section A / Oddelek A

Model of declaration / Vzorec izjave

I, the undersigned/ Spodaj podpisani,

.....
[owner or the natural person who has authorisation in writing from the owner to carry out the non-commercial movement on behalf of the owner⁽¹⁾ / lastnik ali fizična oseba, ki ima pisno dovoljenje lastnika, da opravlja netrgovski premik v imenu lastnika⁽¹⁾]

declare that the following pet animals are not subject to a movement that aims at their sale or a transfer of ownership and will accompany the owner or the natural person who has authorisation in writing from the owner to carry out the non-commercial movement on behalf of the owner⁽¹⁾ within not more than 5 days of his movement. / izjavljam, da cilj premika naslednjih hišnih živali ni njihova prodaja ali prenos lastništva ter da navedene živali spremljajo lastnika ali fizično osebo, ki ima pisno dovoljenje lastnika, da opravlja netrgovski premik v imenu lastnika⁽¹⁾ v največ 5 dneh njegovega premika.

Transponder/tattoo ⁽¹⁾ alphanumeric code / Črkovno-številčna oznaka transponderja/vtetoviranega znamenja ⁽¹⁾	Animal health certificate number / Številka veterinarskega spričevala

During the non-commercial movement, the above animals will remain under the responsibility of / Med netrgovskim premikom je za zgoraj navedene živali odgovorna naslednja oseba:

⁽¹⁾either/bodisi [the owner] / [lastnik]

⁽¹⁾or/bodisi [the natural person who has authorisation in writing from the owner to carry out the non-commercial movement on behalf of the owner] / [fizična oseba, ki ima pisno dovoljenje lastnika, da opravlja netrgovski premik v imenu lastnika].

⁽¹⁾or/bodisi [the natural person designated by the carrier contracted to carry out the non-commercial movement on behalf of the owner / fizična oseba, ki jo določi prevoznik, s katerim je sklenjena pogodba za opravljanje netrgovskega premika živali v imenu lastnika (*insert name of the carrier / vstaviti ime prevoznika*) :]

Place and date: / Kraj in datum:

Signature of the owner or natural person who has authorisation in writing from the owner to carry out the non-commercial movement on behalf of the owner⁽¹⁾ / Podpis lastnika ali fizične osebe, ki ima pisno dovoljenje lastnika, da opravlja netrgovski premik v imenu lastnika⁽¹⁾:

.....
(1) delete as appropriate. / neustrezno črtati.