
Status: Point in time view as at 31/12/2020.

Changes to legislation: There are outstanding changes not yet made to Regulation (EU) No 576/2013 of the European Parliament and of the Council. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

ANNEX IV

Validity requirements for the rabies antibody titration test

1. The collection of the sample of blood necessary to carry out the rabies antibody titration test must be carried out and documented by an authorised veterinarian in the appropriate section of the identification document;
2. The rabies antibody titration test:
 - (a) must be carried out on a sample collected at least 30 days after the date of vaccination and:
 - (i) not less than three months before the date of:
 - the non-commercial movement from a territory or a third country other than those listed in the implementing acts adopted pursuant to Article 13(1) or (2), or
 - the transit through such a territory or third country, where the conditions laid down in point (c) of Article 12 are not fulfilled, or
 - (ii) before the pet animal left the Union for movement to or transit through a territory or a third country other than those listed pursuant to Article 13(1) or (2); the identification document in the format provided for in Article 21(1) must confirm that a rabies antibody titration test was carried out with a favourable result before the date of movement;
 - (b) must measure a level of neutralising antibody to rabies virus in serum equal to or greater than 0,5 IU/ml and using a method prescribed in the relevant part of the Chapter concerning rabies in the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the World Organisation for Animal Health;
 - (c) must be performed in a laboratory approved in accordance with Article 3 of Decision 2000/258/EC;
 - (d) does not have to be renewed following a satisfactory result described in point (b), provided that the pet animal is revaccinated within the period of validity referred to in point 2(e) of Annex III of the previous vaccination.

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