Regulation (EU) 2016/429 of the European Parliament and of the Council of 9 March 2016 on transmissible animal diseases and amending and repealing certain acts in the area of animal health ('Animal Health Law') (Text with EEA relevance)

PART IV

REGISTRATION, APPROVAL, TRACEABILITY AND MOVEMENTS

TITLE I

TERRESTRIAL ANIMALS, GERMINAL PRODUCTS AND PRODUCTS OF ANIMAL ORIGIN FROM TERRESTRIAL ANIMALS

CHAPTER 1

Registration, approval, record-keeping and registers

Section 4

Record-keeping

Article 103

Record-keeping obligations of germinal product establishments

1 Operators of germinal product establishments shall keep and maintain records containing at least the following information:

- a the breed, age, identification and health status of donor animals used for the production of germinal products;
- b the time and place of collection, and the processing and storage, of germinal products collected, produced or processed;
- c the identification of the germinal products together with details of their place of destination, if known;
- d the documents required to accompany germinal products arriving at or leaving the establishment in question in accordance with Article 162 and Article 164(2) and any rules adopted pursuant to Article 162(3) and (4);
- e where relevant, the results of clinical and laboratory tests;
- f laboratory techniques used.

2 Establishments presenting a low risk of spreading listed or emerging diseases may be exempted by the Member State concerned from the requirement to keep records of all or some of the information listed in paragraph 1.

3 Operators of germinal product establishments shall keep the records provided for in paragraphs 1 and 2 on their establishment and:

a make them immediately available to the competent authority on request;

Changes to legislation: There are outstanding changes not yet made to Regulation (EU) 2016/429 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) View outstanding changes

b retain them for a minimum period to be prescribed by the competent authority, which may not be less than three years.

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Changes and effects yet to be applied to the whole legislation item and associated provisions

- Art. 17(1A) words substituted by S.I. 2021/1273 reg. 8Sch. 2 para. (t)