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COVER NOTE

From:	Secretary-General of the European Commission, signed by Ms Martine DEPREZ, Director
date of receipt:	29 January 2021
To:	Mr Jeppe TRANHOLM-MIKKELSEN, Secretary-General of the Council of the European Union
No. Cion doc.:	C(2021) 436 final ANNEXES 1 to 2
Subject:	ANNEXES to the COMMISSION DELEGATED REGULATION supplementing Regulation (EU) 2019/6 of the European Parliament and of the Council as regards the content and format of the information necessary to apply Articles 112(4) and 115(5) and to be contained in the single lifetime identification document referred to in Article 8(4) of that Regulation

Delegations will find attached document C(2021) 436 final ANNEXES 1 to 2.

Encl.: C(2021) 436 final ANNEXES 1 to 2

EUROPEAN
COMMISSION

Brussels, 29.1.2021
C(2021) 436 final

ANNEXES 1 to 2

ANNEXES

to the

COMMISSION DELEGATED REGULATION

supplementing Regulation (EU) 2019/6 of the European Parliament and of the Council as regards the content and format of the information necessary to apply Articles 112(4) and 115(5) and to be contained in the single lifetime identification document referred to in Article 8(4) of that Regulation

ANNEX I

1. The content of the information necessary to apply Article 112(4) of Regulation (EU) 2019/6 shall be the following:

   (a) contact details of the signing veterinarian responsible who treated the equine animal concerned with a veterinary medicinal product authorised under the exemption provided for in Article 8(4) or administered in accordance with Article 112(4) of Regulation (EU) 2019/6;

   (b) the declaration for the equine animal concerned that it is not intended for slaughter for human consumption to be done by the veterinarian responsible in consent with the owner or operator of the equine animal.

2. The content of the information necessary to apply Article 115(5) of Regulation (EU) 2019/6 shall be the following:

   (a) contact details of the signing veterinarian responsible who administered a veterinary medicinal product containing a substance included in the list established in accordance with Article 115(5) of Regulation (EU) 2019/6;

   (b) date and place of the last administration of the veterinary medicinal product referred to in point (a) to the equine animal concerned;

   (c) details of the substance referred to in point (a).

ANNEX II

1. The information necessary to apply Articles 112(4) and 115(5) of Regulation (EU) 2019/6 shall be included in a dedicated section that:

   (a) shall be indivisibly integrated in the single lifetime identification document;

   (b) shall contain titled form fields to be completed in accordance with detailed instructions; those titled form fields and the instructions for their completion shall be displayed in French, English and the official language of the Member State in which the single lifetime identification document is issued;

   (c) shall consist of at least two parts providing form fields for the entry of information necessary:

      (i) to declare the equine animal as not intended for slaughter for human consumption in order to apply Article 112(4);

      (ii) to document the date of last administration of a veterinary medicinal product containing a substance included in the list established in accordance with Article 115(5) of Regulation (EU) 2019/6, and details of that substance.

2. The format of the information necessary to apply Article 112(4) of Regulation (EU) 2019/6 shall meet the following additional criteria:

   (a) the format of the dedicated section referred to in paragraph 1 shall ensure that at least the declaration on the exclusion from slaughter for human consumption can be protected from fraudulent alterations;

   (b) the format of the declaration referred to in point (a) shall be compatible with a corresponding entry in the database referred to in Article 109(1)(d) of Regulation (EU) 2016/429.