Status: EU Directives are being published on this site to aid cross referencing from UK legislation. After IP completion day (31 December 2020 11pm) no further amendments will be applied to this version.

[^{F1}[^{F2}[^{F3}ANNEX C

BRUCELLOSIS

Textual Amendments

- **F1** Substituted by Council Directive 97/12/EC of 17 March 1997 amending and updating Directive 64/432/ EEC on health problems affecting intra-Community trade in bovine animals and swine.
- **F2** Substituted by Commission Regulation (EC) No 535/2002 of 21 March 2002 amending Annex C to Council Directive 64/432/EEC and amending Decision 2000/330/EC.
- F3 Substituted by Commission Decision of 10 December 2008 amending Annex C to Council Directive 64/432/EEC and Decision 2004/226/EC as regards diagnostic tests for bovine brucellosis (notified under document number C(2008) 7642) (Text with EEA relevance) (2008/984/EC).

2. IMMUNOLOGICAL TESTS

- 2.2. Enzyme-linked immunosorbent assays (ELISAs) or other binding assays for the detection of bovine brucellosis in serum or milk
- 2.2.2. Standardisation of the test
- 2.2.2.1. Standardisation of the test procedure for individual serum samples:
- (a) a 1/150 pre-dilution⁽¹⁾ of the OIEISS or a 1/2 pre-dilution of the OIEELISA_{WP}SS or a 1/16 pre-dilution of the OIEELISA_{SP}SS made up in a negative serum (or in a negative pool of sera) must give a positive reaction;
- (b) a 1/600 pre-dilution of the OIEISS or a 1/8 pre-dilution of the OIEELISA_{WP}SS or a 1/64 pre-dilution of the OIEELISA_{SP}SS made up in a negative serum (or in a negative pool of sera) must give a negative reaction;
- (c) the OIEELISA_NSS must always give a negative reaction.]]]

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(1) [^{F1}[^{F2}[^{F3}For the purpose of this Annex, dilutions given for making up liquid reagents are expressed as, for example, 1/150 shall mean a 1 in 150 dilution.]]]

Textual Amendments

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