

*Status: Point in time view as at 03/02/2016.*

*Changes to legislation: There are currently no known outstanding effects for the Regulation (EC) No 999/2001 of the European Parliament and of the Council, CHAPTER B. (See end of Document for details)*

## [<sup>F1</sup>ANNEX III

### MONITORING SYSTEM

#### Textual Amendments

- F1** Substituted by [Commission Regulation \(EC\) No 2245/2003 of 19 December 2003 amending Annex III to Regulation \(EC\) No 999/2001 of the European Parliament and of the Council as regards monitoring of transmissible spongiform encephalopathies in ovine and caprine animals.](#)

## [<sup>F2</sup>CHAPTER B

### REPORTING AND RECORDING REQUIREMENTS

- I. REQUIREMENTS ON MEMBER STATES
- A. **Information to be presented by Member States in their annual report as provided for in Article 6(4)**
1. The number of suspected cases placed under official movement restrictions in accordance with Article 12(1), per animal species.
  2. The number of suspected cases subject to laboratory examination in accordance with Article 12(2), per animal species, including the results of the rapid and confirmatory tests (number of positives and negatives) and, with regard to bovine animals, the age distribution of all tested animals. The age distribution should be grouped as follows: 'below 24 months', distribution per 12 months between 24 and 155 months, and 'above 155 months' of age.
  3. The number of flocks where suspected cases in ovine and caprine animals have been reported and investigated pursuant to Article 12(1) and (2).
  4. The number of bovine animals tested within each subpopulation referred to in Chapter A, Part I, points 2.1, 2.2, 3.1 and 5. The method of the sample selection, the results of the rapid and confirmatory tests and the age distribution of the tested animals grouped as set out in point 2 shall be provided.
  5. The number of ovine and caprine animals and flocks tested within each subpopulation referred to in Chapter A, Part II, points 2, 3, 5 and 6 together with the method for sample selection and the results of the rapid and confirmatory tests.
  6. The geographical distribution, including the country of origin if not the same as the reporting country, of positive cases of BSE and scrapie. The year, and where possible the month of birth shall be given for each TSE case in bovine, ovine and caprine animals. TSE cases which have been considered atypical shall be indicated. For scrapie cases, the results of the primary and secondary molecular testing, referred to in Annex X, Chapter C, point 3.2(c), shall be reported, where appropriate.
  7. In animals other than bovine, ovine and caprine animals, the number of samples and confirmed TSE cases per species.
  8. The genotype, and, where possible, the breed, of each ovine animal either found positive to TSE and sampled in accordance with Chapter A, Part II, point 8.1, or sampled in accordance with Chapter A, Part II, point 8.2.

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## B. Reporting periods

The compilation of reports containing the information referred to in Section A and submitted to the Commission (which shall send it to the European Food Safety Authority) on a monthly basis in the electronic format agreed between the Member States, the Commission and the European Food Safety Authority or, with regard to the information referred to in point 8 on a quarterly basis, may constitute the annual report as required by Article 6(4), provided that the information is updated whenever additional information becomes available.

### II. INFORMATION TO BE PRESENTED IN THE UNION SUMMARY REPORT

The Union summary shall be presented in a tabled format covering at least the information referred to in Part I.A for each Member State.

From 1 January 2016, the European Food Safety Authority shall analyse the information referred to in Part I and publish by the end of November a summary report on the trends and sources of Transmissible Spongiform Encephalopathies in the Union.

### III. RECORDS

1. The competent authority shall keep, for 7 years, records of the information referred to in Part I.A.
2. The investigating laboratory shall keep, for 7 years, all records of testing, in particular laboratory workbooks and, where appropriate, paraffin blocks and photographs of western blots.]]

#### Textual Amendments

- F2** Substituted by [Commission Regulation \(EU\) 2016/27 of 13 January 2016 amending Annexes III and IV to Regulation \(EC\) No 999/2001 of the European Parliament and of the Council laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies.](#)

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