Council Directive of 26 March 1990 laying down the conditions governing the preparation, placing on the market and use of medicated feedingstuffs in the Community (90/167/EEC)

## Article 10

- 1 Member States shall ensure that, without prejudice to animal-health rules, there are no prohibitions, limitations or obstacles in respect of intra-Community trade
- in medicated feedingstuffs which have been manufactured in accordance with the requirements of this Directive, and in particular Article 4 thereof, with authorized premixes which have the same active substances as pre-mixes authorized by the Member State of destination, in accordance with the criteria of Directive 81/852/EEC, and a quantitative and qualitative composition similar thereto,
- subject to the specific provisions of Council Directive 86/469/EEC of 16 September 1986 concerning the examination of animals and fresh meat for the presence of residues<sup>(1)</sup> and Council Directive 88/299/EEC of 17 May 1988 on trade in animals treated with certain substances having a hormonal action and their meat, as referred to in Article 7 of Directive 88/146/EEC<sup>(2)</sup>, in animals to which those medicated feedingstuffs except those produced pursuant to the second subparagraph of Article 3 (1), have been administered, or in meat, flesh, offal or their products from such animals.
- Where the application of paragraph 1 gives rise to dispute, in particular as concerns recognition of the similar nature of the pre-mix, the Member States concerned or the Commission may submit the dispute to assessment by an expert appearing on a list of Community experts to be drawn up by the Commission on a proposal from the Member States.

If the two Member States so agree beforehand, the parties shall abide by the opinion of the expert, in compliance with Community legislation.

3 The Member State of destination may require that each consignment of a medicated feedingstuff be accompanied by a certificate issued by the competent authority, corresponding to the specimen form in Annex B.

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.

- (1) OJ No L 275, 26. 9. 1986, p. 36.
- (2) OJ No L 128, 21. 5. 1988, p. 36.