

## LIST OF LEGISLATION

- Council Directive 65/65/EEC of 26 January 1965 on the approximation of provisions laid down by law, regulation or administrative action relating to proprietary medicinal Official Journal No L 022, 09/02/1965 P. 0369 – 0373
- Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances Official Journal No L 196, 16/08/1967 P. 0001 – 0005
- Council Directive 78/25/EEC of 12 December 1977 on the approximation of the laws of the Member States relating to the colouring matters which may be added to medicinal products Official Journal No L 011, 14/01/1978 P. 0018 – 0020
- Council Directive 81/851/EEC of 28 September 1981 on the approximation of the laws of the Member States relating to veterinary medicinal products Official Journal No L 317, 06/11/1981 P. 0001 – 0015
- Council Directive 81/852/EEC of 28 September 1981 on the approximation of the laws of the Member States relating to analytical, pharmaco-toxicological and clinical standards and protocols in respect of the testing of veterinary medicinal products Official Journal No L 317, 06/11/1981 P. 0016 – 0028
- Commission Directive 84/449/EEC of 25 April 1984 adapting to technical progress for the sixth time Council Directive 67/548/EEC on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances
- Council Directive 87/18/EEC of 18 December 1986 on the harmonisation of laws, regulations and administrative provisions relating to the application of the principles of good laboratory practice and the verification of their applications for tests on chemical substances Official Journal No L 015, 17/01/1987 P. 0029 – 0030
- Commission Directive 87/302/EEC of 18 November 1987 adapting to technical progress for the ninth time Council Directive 67/548/EEC on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances
- 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 Official Journal No L 128, 21/05/1988 P. 0036 – 0038
- Council Directive 88/320/EEC of 9 June 1988 on the inspection and verification of Good Laboratory Practice (GLP) Official Journal No L 145, 11/06/1988 P. 0035 – 0037
- Council Directive 90/167/EEC of 26 March 1990 laying down the conditions governing the preparation, placing on the market and use of medicated feedingstuffs in the Official Journal No L 092, 07/04/1990 P. 0042 – 0048
- Council Directive 90/676/EEC of 13 December 1990 amending Directive 81/851/EEC on the approximation of the laws of the Member States relating to veterinary medicinal products Official Journal No L 373, 31/12/1990 P. 0015 – 0025

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- Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market Official Journal No L 230, 19/08/1991 P. 0001 – 0032
- Commission Directive 91/632/EEC of 28 October 1991 adapting to technical progress for the fifteenth time Council Directive 67/548/EEC on the approximation of the laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances Official Journal No L 338, 10/12/1991 P. 0023
- Council Directive 90/220/EEC of 23 April 1990 on the deliberate release into the environment of genetically modified organisms Official Journal No L 117, 08/05/1990 P. 0015 – 0027
- Council Regulation (EEC) No 2377/90 of 26 June 1990 laying down a Community procedure for the establishment of maximum residue limits of veterinary medicinal products in foodstuffs of animal origin Official Journal No L 224, 18/08/1990 P. 0001 – 0008
- Council Directive 90/677/EEC of 13 December 1990 extending the scope of Directive 81/851/EEC on the approximation of laws of the Member States relating to veterinary medicinal products and laying down additional provisions for immunological veterinary medicinal products Official Journal No L 373 of 31.12.1990
- Council Directive 91/412/EEC Good Manufacturing Practice for veterinary medicinal products Official Journal No L 228 of 17.8.1991
- Commission Directive 92/18/EEC of 20 March 1992 modifying the Annex to Council Directive 81/852/EEC on the approximation of the laws of Member States relating to analytical, pharmacotoxicological and clinical standards and protocols in respect of the texturing of veterinary medicinal products Official Journal No L 097 , 10/04/1992 P. 0001 - 0023
- Council Directive 92/74/EEC of 22 September 1992 widening the scope of Directive 81/851/EEC on the approximation of laws of the Member States relating to veterinary medicinal products and laying down additional provisions for homeopathic veterinary medicinal products Official Journal No L 297 of 13.10.1992
- Council Regulation 2309/93 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Agency for the Evaluation of Medicinal Products Official Journal No L 214, 24.08.1993
- Commission Directive 93/67/EEC of 20 July 1993 laying down the principles for assessment of risks to man and the environment of substances notified in accordance with Council Directive 67/548/EEC Official Journal No L 227, 08/09/1993 P. 0009 – 0018
- Commission Regulation (EC) No 1488/94 of 28 June 1994 laying down the principles for the assessment of risks to man and the environment of existing substances in accordance with Council Regulation (EEC) No 793/93 (Text with EEA relevance) Official Journal No L 161, 29/06/1994 P. 0003
- Commission Regulation (EC) No 541/95 of 10 March 1995 concerning the examination of variations to the terms of a marketing authorization granted by a competent authority of a Member State Official Journal No L 055 , 11/03/1995 P. 0007 – 0014

- Commission Regulation (EC) No 542/95 of 10 March 1995 concerning the examination of variations to the terms of a marketing authorization falling within the scope of Council Regulation (EEC) No 2309/93 Official Journal No L 055 , 11/03/1995 P. 0015 - 0021
- Directive 96/23/EEC on measures to monitor certain substances and residues thereof in live animals and animal products and repealing Directives 85/358/EEC and 86/469/EEC and Decisions 89/187/EEC and 91/664/EEC Official Journal No L 125, 23.5.1996, p. 10