

Council Regulation (EC) No 297/95 of 10 February 1995 on fees payable to the European Agency for the Evaluation of Medicinal Products

Article 1	Scope
Article 2	The Agency shall indicate in its annual estimate intended for...
Article 3	Medicinal products for human use covered by the procedures laid down in Regulation (EC) No 726/2004
Article 4	Medicinal products for human use covered by the procedures laid down in Directive 2001/83/EC
Article 5	Medicinal products for veterinary use covered by the procedures laid down in Regulation (EC) No 726/2004
Article 6	Veterinary medicinal products covered by the procedures laid down in Directive 2001/82/EC
Article 7	Establishment of maximum residue limits (MRL) for veterinary medicinal products in accordance with the procedures laid down in Regulation (EEC) No 2377/90
Article 8	Various Fees
Article 9	Possible fee reductions
Article 10	Due date and deferral of the payment
Article 11	Implementing rules
Article 12	Amendment
Article 13	Entry into force and legal effect
	Signature

Changes to legislation: There are outstanding changes not yet made to Council Regulation (EC) No 297/95. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) [View outstanding changes](#)

- (1) [^{X1}Opinion delivered on 19 January 1995 (not yet published in the Official Journal).]
- (2) [OJ No L 214, 24.8.1993, p. 1.](#)
- (3) [OJ No 22, 9.2.1965, p. 369/65.](#) Directive as last amended by Directive 93/39/EEC ([OJ No L 214, 24.8.1993, p. 22.](#))
- (4) [OJ No L 317, 6.11.1981, p. 1.](#) Directive as last amended by Directive 93/40/EEC ([OJ No L 214, 24.8.1993, p. 31.](#))

Editorial Information

X1 Inserted by [Corrigendum, OJ No L 75, 4.4.1995, p. 29 \(297/95\).](#)

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Changes and effects yet to be applied to :

- Regulation revoked in part by [S.I. 2019/775 Sch. 9 para. 1\(b\)](#)