

Commission Regulation (EC) No 668/2009 of 24 July 2009 implementing Regulation (EC) No 1394/2007 of the European Parliament and of the Council with regard to the evaluation and certification of quality and non-clinical data relating to advanced therapy medicinal products developed by micro, small and medium-sized enterprises (Text with EEA relevance)

Article 1	Scope
Article 2	Procedure for evaluation and certification
Article 3	Site visits
Article 4	Combined advanced therapy medicinal products
Article 5	Scientific guidelines
Article 6	Report
Article 7	This Regulation shall enter into force on the 20th day... Signature

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EC) No 668/2009. 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) [OJ L 324, 10.12.2007, p. 121.](#)
- (2) [OJ L 124, 20.5.2003, p. 36.](#)
- (3) [OJ L 311, 28.11.2001, p. 67.](#)
- (4) [OJ L 169, 12.7.1993, p. 1.](#)
- (5) [OJ L 189, 20.7.1990, p. 17.](#)

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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\(q\)](#)