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)

(**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.

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.
View outstanding changes

Changes and effects yet to be applied to :

– Regulation revoked in part by S.I. 2019/775 Sch. 9 para. 1(q)