

Commission Regulation (EC) No 507/2006 of 29 March 2006 on the conditional marketing authorisation for medicinal products for human use falling within the scope of Regulation (EC) No 726/2004 of the European Parliament and of the Council (Text with EEA relevance)

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Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EC) No 507/2006. 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 136, 30.4.2004, p. 1.](#)
- (2) [OJ L 311, 28.11.2001, p. 67.](#) Directive as last amended by Directive 2004/27/EC ([OJ L 136, 30.4.2004, p. 34.](#))
- (3) [OJ L 268, 3.10.1998, p. 1.](#) Decision as last amended by Regulation (EC) No 1882/2003 ([OJ L 284, 31.10.2003, p. 1.](#))
- (4) [OJ L 18, 22.1.2000, p. 1.](#)

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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\(k\)](#)