Regulation (EC) No 141/2000 of the European Parliament and of the Council of 16 December 1999 on orphan medicinal products

Article 1	Purpose
Article 2	Definitions
11111111	
Article 3	Criteria for designation
Article 4	Committee for Orphan Medicinal Products
Article 5	Procedure for designation and removal from the register
Article 6	Protocol assistance
Article 7	Community marketing authorisation
Article 8	Market exclusivity
Article 9	Other incentives
Article 10	General report
Article 10a	(1) The Commission shall be assisted by the Standing
	Committee
Article 10b	Exercise of the delegation
Article 11	Entry into force
	Signature

Changes to legislation: There are outstanding changes not yet made to Regulation (EC) No 141/2000 of the European Parliament and of the Council. 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 C 276, 4.9.1998, p. 7.
- (2) OJ C 101, 12.4.1999, p. 37.
- (3) Opinion of the European Parliament of 9 March 1999 (OJ C 175, 21.6.1999, p. 61), Council Common Position of 27 September 1999 (OJ C 317, 4.11.1999, p. 34) and Decision of the European Parliament of 15 December 1999 (not yet published in the Official Journal).
- (4) OJ 22, 9.2.1965, p. 369. Directive as last amended by Directive 93/39/EEC (OJ L 214, 24.8.1993, p. 22).
- (**5**) OJ L 155, 22.6.1999, 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(f)