

Council Directive 93/41/EEC of 14 June 1993 repealing Directive 87/22/EEC on the approximation of national measures relating to the placing on the market of high-technology medicinal products, particularly those derived from biotechnology

- Article 1 With effect from 1 January 1995, Directive 87/22/EEC is hereby...
- Article 2 Applications for marketing authorizations which have been referred to the...
- Article 3 Member States shall take all appropriate measures to comply with...
- Article 4 This Directive is addressed to the Member States.
Signature

Status: EU Directives are being published on this site to aid cross referencing from UK legislation. After IP completion day (31 December 2020 11pm) no further amendments will be applied to this version.

- (1) [OJ No C 58, 8. 3. 1990, p. 1.](#)
- (2) [OJ No C 183, 15. 7. 1991, p. 145](#) and [OJ No C 150, 31. 5. 1993.](#)
- (3) [OJ No C 269, 14. 10. 1991, p. 84.](#)
- (4) [OJ No L 15, 17. 1. 1987, p. 38.](#)
- (5) See page 1 of this Official Journal.
- (6) [OJ No L 81, 26. 3. 1988, p. 75.](#)
- (7) See page 22 of this Official Journal.
- (8) See page 31 of this Official Journal.