Regulation (EC) No 1901/2006 of the European Parliament and of the Council of 12 December 2006 on medicinal products for paediatric use and amending Regulation (EEC) No 1768/92, Directive 2001/20/EC, Directive 2001/83/EC and Regulation (EC) No 726/2004 (Text with EEA relevance) (revoked)

TITLE IV

POST-AUTHORISATION REQUIREMENTS

Article 33

Textual Amendments applied to the whole legislation

F1 Regulation revoked (31.12.2020) by The Human Medicines (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/775), Sch. 9 para. 1(1) (subject to transitional provisions in S.I. 2012/1916, Sch. 33A); 2020 c. 1, Sch. 5 para. 1(1)

Changes to legislation:

This version of this Regulation was derived from EUR-Lex on IP completion day (31 December 2020 11:00 p.m.). It has not been amended by the UK since then. Find out more about legislation originating from the EU as published on legislation.gov.uk.