

Commission Regulation (EC) No 1234/2008 of 24 November 2008 concerning the examination of variations to the terms of marketing authorisations for medicinal products for human use and veterinary medicinal products (Text with EEA relevance) (revoked)

CHAPTER II

VARIATIONS TO MARKETING AUTHORISATIONS GRANTED IN ACCORDANCE WITH DIRECTIVE 87/22/EEC, CHAPTER 4 OF DIRECTIVE 2001/82/EC OR CHAPTER 4 OF DIRECTIVE 2001/83/EC

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Textual Amendments applied to the whole legislation

F1 Regulation revoked (17.5.2024) by [The Veterinary Medicines \(Amendment etc.\) Regulations 2024 \(S.I. 2024/567\)](#), regs. 1(1), **209**

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](https://www.legislation.gov.uk).