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**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://legislation.gov.uk). (See end of Document for details)*

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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 III **U.K.**

**VARIATIONS TO CENTRALISED MARKETING AUTHORISATIONS**

*Article 15* **U.K.**

**Notification procedure for minor variations of type IB**

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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**

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