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

. . . . . .

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**