[F1SCHEDULE 1B

Qualifying Northern Ireland good (QNIG) certificates

Textual Amendments

F1 Sch. 1B inserted (E.W.S.) (31.12.2020) by The Veterinary Medicines and Residues (Amendment) (EU Exit) Regulations 2020 (S.I. 2020/1461), regs. 1(2)(b), **4(8)**

4. The condition is that the person who holds a marketing authorisation in respect of the product which is valid in Northern Ireland under the Northern Ireland VMRs, who must be a person established in Northern Ireland, has provided the Secretary of State with the following information—

(a) the Northern Ireland address of that person;

(b) all necessary administrative information, and all scientific documentation necessary for demonstrating the safety, quality and efficacy of the veterinary medicinal product, equivalent to that which would need to be provided under Schedule 1 if an application for a marketing authorisation was to be made in respect of that product under paragraph 1 of that Schedule (allowing for any relevant derogations provided for in Part 2 of that Schedule);

(c) the name and address of a person who resides in the United Kingdom or in a member State who is to provide in respect of the veterinary medicinal product, permanently and continuously, the services of a qualified person (pharmacovigilance) for the purposes of Part 8 of Schedule 1.**]**

Changes to legislation:
There are currently no known outstanding effects for the The Veterinary Medicines Regulations 2013, Paragraph 4.