VPA10846/018/001

INMEVA, suspension for injection

Variation	Summary	Date
Vet - B42	VNRA - Vet - B42 - - Vet - B42- Implementation in practice of changes already foreseen in an approved change management protocol (CMP) for the finished product	31/07/26
Vet - B42	VNRA - Vet - B42 - Implementation in practice of changes already foreseen in an approved change management protocol (CMP) for the finished product - B42 Changes to the quality part of the dossier: Implementation in practice of changes already foreseen in an approved change management protocol (CMP) for the finished product	17/02/25
Vet - G.I.18	VRA-S - Vet - G.I.18 - One-off alignment of the product information with version 9.0 (or the latest version of the QRD templates that are in effect at the time that this one-off variation is submitted) of the QRD templates i.e. major update of the QRD templates in accordance with Regulation (EU) 2019/6, for veterinary medicinal products placed on the market in accordance with Directive 2001/82/EC or Regulation (EC) No 726/2004 - G.I.18 Safety, Efficacy, Pharmacovigilance changes - One-off alignment of the product information with version 9.0 (or the latest version of the QRD templates that are in effect at the time that this one-off variation is submitted) of the QRD templates i.e. major update of the QRD templates in accordance with Regulation (EU) 2019/6, for veterinary medicinal products placed on the market in accordance with Directive 2001/82/EC or Regulation (EC) No 726/2004	12/12/24
Vet - B42	VNRA - Vet - B42 - Implementation in practice of changes already foreseen in an approved change management protocol (CMP) for the finished product - B42 Changes to the quality part of the dossier: Implementation in practice of changes already foreseen in an approved change management protocol (CMP) for the finished product	03/04/24
B.II.d.2.d	IB - B.II.d.2.d - d) Other changes to a test procedure (including replacement or addition) - B.II.d.2.d - QUALITY CHANGES - FINISHED PRODUCT - Control of finished product - Change in test procedure for the finished product - Other changes to a test procedure (including replacement or addition)	28/02/22
B.II.f.1.b.5	IB - B.II.f.1.b.5 - 5. Extension of the shelf-life of a biological/ immunological medicinal product in accordance with an approved stability protocol - B.II.f.1.b.5 - QUALITY CHANGES - FINISHED PRODUCT - Stability - Change in the shelf-life or storage conditions of the finished product - Extension of the shelf life of the finished product - Extension of the shelf-life of a biological/ immunological medicinal product in accordance with an approved stability protocol	10/02/22