

VPA10454/068/001

**Ivomec Super Injection for Cattle**

<b>Variation</b>	<b>Summary</b>	<b>Date</b>
Vet - F.II.b.3 g)	VRA-S - Vet - F.II.b.3 g) - g) Move the sterilizing filtration from A/B to C - F.II.b.3 g) Quality Changes - Finished Product -Manufacture - Change in the manufacturing process of the finished product, including an intermediate used in themanufacture of the finished product - Move the sterilizing filtration from A/B to C	21/12/23
B.I.a.1.b	II - B.I.a.1.b - b) Introduction of a manufacturer of the active substance supported by an ASMF - B.I.a.1.b - QUALITY CHANGES - ACTIVE SUBSTANCE - Manufacture - Change in the manufacturer of a starting material/reagent/intermediate used in the manufacturing process of the active substance or change in the manufacturer (including where relevant quality control testing sites) of the active substance, where no Ph. Eur. Certificate of Suitability is part of the approved dossier - Introduction of a manufacturer of the active substance supported by an ASMF	18/05/23
Vet - C1	VNRA - Vet - C1 - Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV) - C1 Changes to the safety, efficacy and pharmacovigilance part of the dossier: Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV)	15/03/23
Vet - A1 e)	VNRA - Vet - A1 e) - e) Change in the name or address or contact details of a manufacturer or importer of the finished product (including batch release or quality control testing sites) - A1 e) Administrative changes: Change in the name or address or contact details of a manufacturer or importer of the finished product (including batch release or quality control testing sites)	04/08/22