VPA10846/015/001

Previron 200 mg/ml solution for injection for pigs

Variation	Summary	Date
Vet - A1 b)	VNRA - Vet - A1 b) - b) Change in the name or address or contact details of a manufacturer or supplier of the active substance, starting material, reagent or intermediate used in the manufacture of the active substance or a quality control testing site (where ecified in the dossier) where no European Pharmacopoeia (Ph. Eur.) Certificate of Suitability (CEP) is part of the approved dossier. - A1 b) Administratvie changes: Change in the name or address or contact details of a manufacturer or supplier of the active substance, starting material, reagent or intermediate used in the manufacture of the active substance or a quality control testing site (where ecified in the dossier) where no European Pharmacopoeia (Ph. Eur.) Certificate of Suitability (CEP) is part of the approved dossier.	08/02/24
Vet - F.I.c.1 b)	VRA-R - Vet - F.I.c.1 b) - b) Liquid active substances (non sterile) - F.I.c.1 b) Quality Changes - Active Substance - Container closure system - Change in immediate packaging of the active substance - Liquid active substances (non sterile)	08/02/24
Vet - F.I.c.2.z	VRA-R - Vet - F.I.c.2.z - z) Other changes under this code level e.g. variations outlined in section 6 and 7 of EMA/CMDv/7381/2021 - F.I.c.2 z) Quality Changes - Active Substance - Container closure system - Change in the specification parameters and/or limits of the immediate packaging of the active substance - Other changes under this code level, e.g. variations outlined in section 6 and 7 of EMA/CMDv/7381/2021	08/02/24