VPA10990/054/001

Redymox 150 mg/ml suspension for injection for cattle, sheep, pigs, dogs and cats

Variation	Summary	Date
Vet - F.I.d.1 c)	VRA-R - Vet - F.I.d.1 c) - - Vet - F.I.d.1 c) - Change in the re-test period/storage period of the active substance where no Ph. Eur. Certificate of Suitability covering the retest period is part of the approved dossier - Extension or introduction of a re-test period/storage period supported by real time data	21/10/26
Vet - F.III.1 a) 1.	VRA-R - Vet - F.III.1 a) 1. - - Vet - F.III.1 a) 1. - Submission of a new or updated Ph. Eur. certificate of suitability or deletion of Ph. Eur. certificate of suitability - For an active substance, For a starting material/reagent/intermediate used in the manufacturing process of the active substance, For an excipient - European Pharmacopoeial Certificate of Suitability to the relevant Ph. Eur. Monograph. - New certificate for a non-sterile active substance that is to be used in a sterile medicinal product, where mwater is used in the last steps of the synthesis and the material is not claimed to be endotoxin free	21/10/26
Vet - F.III.1 a) 1.	VRA-R - Vet - F.III.1 a) 1. - - Vet - F.III.1 a) 1. - Submission of a new or updated Ph. Eur. certificate of suitability or deletion of Ph. Eur. certificate of suitability - For an active substance, For a starting material/reagent/intermediate used in the manufacturing process of the active substance, For an excipient - European Pharmacopoeial Certificate of Suitability to the relevant Ph. Eur. Monograph. - New certificate for a non-sterile active substance that is to be used in a sterile medicinal product, where mwater is used in the last steps of the synthesis and the material is not claimed to be endotoxin free	25/06/26
Vet - F.I.d.1 c)	VRA-R - Vet - F.I.d.1 c) - - Vet - F.I.d.1 c) - Change in the re-test period/storage period of the active substance where no Ph. Eur. Certificate of Suitability covering the retest period is part of the approved dossier - Extension or introduction of a re-test period/storage period supported by real time data	24/06/26
Vet - F.III.1 a) 1.	VRA-R - Vet - F.III.1 a) 1. - - Vet - F.III.1 a) 1. - Submission of a new or updated Ph. Eur. certificate of suitability or deletion of Ph. Eur. certificate of suitability - For an active substance, For a starting material/reagent/intermediate used in the manufacturing process of the active substance, For an excipient - European Pharmacopoeial Certificate of Suitability to the relevant Ph. Eur. Monograph. - New certificate for a non-sterile active substance that is to be used in a sterile medicinal product, where mwater is used in the last steps of the synthesis and the material is not claimed to be endotoxin free	24/06/26
Vet - G.I.15 z)	VRA-R - Vet - G.I.15 z) - - Vet - G.I.15 z) - Changes to the labelling or the package leaflet which are not connected with the summary of product characteristics - Other changes under this code level, e.g. variations outlined in section 6 and 7 of this	29/08/25

	guidance	