Veterinary Medicines Directorate

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Post Authorisation Assessments

Doxx-Sol 500 mg/g Powder for Use in Drinking Water/Milk Replacer for Pre-ruminant Calves, Pigs and Chickens

Vm 30282/4022

17 July 2026	Change in the address of the local representative to Huvepharma Ltd, 2nd Floor South, One Castle Park, Tower Hill, Bristol, BS2 0JA, United Kingdom Tel: +32 3 318 36 54. (GB + NI).
12 June 2026	Submission of an updated Ph. Eur. Certificate for an active substance. (GB)
5 June 2026	Submission of a Ph. Eur. CEP for an active substance (NI)
18 September 2025	Alignment of the product information with version 9.0* of the QRD templates.
17 July 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI).
11 May 2024	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance. (NI)
28 July 2023	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
22 June 2020	Deletion of manufacturing site for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
17 February 2020	Renewal - UK as CMS.
21 August 2017	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.