



Post Authorisation Assessments

Doxatib 500 mg/g Powder for Use in Drinking Water for Pigs and Chickens Vm 01656/3053

16 May 2026	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance.
08 May 2026	Change in address of a manufacturer of the finished product.
25 September 2025	Change in distributor details to KRKA UK Ltd, Thames House, Waterside Drive, Langley, SL3 6EZ, United Kingdom.
23 January 2024	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
21 July 2023	One-off alignment of the product information with version 9.0* 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.
04 July 2023	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer.
25 February 2022	Updates to labelling.
24 November 2021	Change in the name and/or address of a manufacturer of the finished product.
16 September 2021	Renewal – UK as CMS.
07 January 2021	Change in the SPC, labelling or package leaflet following assessment of the same change for the reference product.
29 May 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
28 January 2020	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
23 October 2018	Update to the Local Representative details.
09 November 2016	Change in Distributor Details