



Post Authorisation Assessments

Doxatib 500 mg/g Powder for Use in Drinking Water for Pigs and Chickens Vm 01656/5053

•	19 December 2023	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.
•	28 April 2023	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
•	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