



## Post Authorisation Assessments

### Amatib 800 mg/g Oral Powder for Pigs and Chickens

Vm 01656/5070

16 September 2025	Change of Distributor from: Unidrug Distribution Group Limited, Amber Park 1, 2, 3 and 5 Berristow Lane, South Normanton, Alfreton, Derbyshire, DE55 2FH; Centaur Services Limited, Centaur House, Torbay Road, Castle Cary, Somerset, BA7 7EU; Movianto UK Limited, 1A Progress Park, Elstow, Bedford, Bedfordshire, MK42 9XE; National Veterinary Services Limited, Unit 4, Jamage Industrial Estate, Talke Pits, Stoke-on-Trent, Staffordshire, ST7 1XW; Potter Logistics Limited, Charleywood Road, Knowsley Industrial Park, Liverpool, L33 7SG; Henry Schein UK Holdings Limited, Medcare North, Centurion Close, Gillingham Business Park, Gillingham, Kent, ME8 0SB To: KRKA UK Ltd, Thames House, Waterside Drive, Langley, SL3 6EZ, United Kingdom.
18 February 2025	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance.
18 June 2024	One-off alignment of the product information with version 2 of the GB QRD templates.
30 January 2023	Updated certificate of suitability from an already approved manufacturer.
12 August 2022	Updated certificate of suitability from an already approved manufacturer.
04 June 2021	Increase in the shelf-life of the finished product as packaged for sale, from 2 years to 3 years.
21 April 2021	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
18 September 2020	Renewal – UK as CMS.
02 September 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
11 June 2019	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
22 February 2018	Change in contact details for local representative.
23 May 2017	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer. Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer. Change in the specification limits of an excipient.
02 February 2017	Change in pack size of the finished product.

•	09 November 2016	Change in dimensions of the container (immediate packaging)
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