



Post Authorisation Assessments

Advocin 180, 180mg/ml, Solution for Injection for Cattle Vm 42058/5107

15 October 2025	Addition of a test procedure for the active substance.
05 February 2025	Deletion of a non-significant specification parameter for an active substance.
06 October 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.
17 January 2023	Deletion of a test procedure for the finished product. Deletion of a manufacturer of the finished product. Deletion of a manufacturer of the finished product.
29 March 2022	Deletion of manufacturing site for an active substance. Extension of a re-test period of the active substance.
15 December 2021	Addition to a test procedure for an excipient.
08 January 2020	Change in the address of the marketing authorisation holder from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP.
08 January 2020	Replacement of a site for the manufacture of the active substance intermediate.
07 November 2018	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
06 September 2018	Addition of a site where batch control/testing takes place. Replacement of a manufacturer responsible for batch release of the finished product. Addition of a secondary packaging site of the finished product. Addition to a test procedure for the finished product. Changes to a test procedure for the finished product. Decrease in batch size range of the finished product. Addition of an in-process control test. Addition of a manufacturing site of the finished product.
24 July 2018	Change in the name of a manufacturer of the finished product, also responsible for batch release.
08 August 2017	Tightening of specification limits of an active substance used in the manufacturing process of the active substance. Changes to a test procedure (including replacement * or addition*) for the active substance. Change in the specification parameters and/or limits of

	the immediate packaging of the active substance. Additional manufacturer of the active substance only for synthesis.
18 July 2017	Change in the name of the manufacturer of the finished product
14 October 2015	To add a site of secondary packaging for the finished product. To add a new batch range for the finished product. To add a site of manufacture and primary packaging for the finished product.
05 June 2015	Change in the QPPV and/or QPPV contact details and/or back-up procedure.
06 March 2015	Updates to the SPC and product literature.
10 December 2014	Change in the name of a site responsible for manufacturing and batch control and release of the finished product.
20 December 2013	Change to test procedures for the finished product.
30 October 2013	Addition of a finished product manufacturer responsible for batch release. Update to the QPPV contact details.
22 October 2013	Change to the name of the MAH in Austria, Belgium, France and Luxembourg only.
16 August 2013	Change in distributor and MAH from Pfizer Limited to Zoetis UK Limited.
13 April 2012	Change to in process tests applied during the manufacture of the finished product.
20 March 2012	Change in product packaging components.
13 February 2012	Change to the name of the MAH in Spain only.
16 June 2011	Change to the name of the MAH in Poland only.
04 March 2010	Change in test procedure on the finished product.
27 April 2009	Change in test procedure on the finished product.
29 December 2008	Addition of a statement on use of fluoroquinolones to the SPC.
12 May 2008	Repeat use procedure.
15 May 2006	Renewal.
09 May 2006	Change of marketing authorisation holder (MAH) in Portugal only.
12 August 2005	Addition of a new therapeutic indication.
07 July 2005	Addition of a distributor.
25 October 2004	Change of name and address of a manufacturer.
06 August 2003	Addition of a new route of administration.
11 July 2002	Change of supplier of an intermediate in the manufacturing process of the active substance.