



Post Authorisation Assessments

Clamoxyl Ready-to-Use 150 mg/ml Suspension for Injection Vm 42058/4016

13 May 2025	Submission of new CEP for the manufacture of an active substance.
08 April 2022	Change(s) in the SPC and Package Leaflet to implement the outcome of a procedure concerning PSUR.
21 December 2021	Submission of a new certificate of suitability for an active substance.
21 August 2020	Change in the address of the MAH, 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.
28 March 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
13 January 2017	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Addition of an alternative sterilisation site for the active substance.
26 June 2014	Change to the Marketing Authorisation Holder and distributor details.
19 December 2011	Changes to an existing pharmacovigilance system as described in the DDPS.
03 December 2009	Renewal.
10 August 2009	Change of withdrawal period from 21 days to 54 days for meat from cattle, and from 48 hours to 60 hours for milk from cattle.
22 January 2009	Addition of a manufacturer of the active substance.
05 November 2008	Change of legal category from POM to POM-V. Changes to the SPC and Product Literature to bring in line with new legislation.
05 August 2005	Change of site for part of the manufacturing process of the active substance.
17 June 2005	Change of distributor.
18 June 2004	Changes to packaging.
27 February 2003	Change of MAH name.
08 March 2002	Minor change in manufacturing process of the finished product.
05 December 1997	Change of manufacturer of the active substance.