Veterinary Medicines Directorate

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Post Authorisation Assessments

Clamoxyl Long Acting 150 mg/ml Suspension for Injection

Vm 42058/4013

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
10 November 2009	Change of withdrawal period, from 21 days to 92 days for meat from cattle. Removal of a route of administration. Change of dosage details.
10 November 2009	Renewal.
22 July 2009	Change of name of manufacturer of the finished product.
22 January 2009	Addition of a manufacturer of the active substance.
28 October 2008	Change of legal category from POM to POM-V. Changes to the SPC and Product Literature to bring in line with new legislation.
28 July 2005	Change of manufacturer involved in part of the manufacturing process of the active substance.
27 June 2005	Change of distributor.
27 February 2003	Change of name of MAH.
08 March 2002	Change to method of sterilisation of container.
23 August 2000	Change of manufacturer of the active substance.
28 January 1998	Change of manufacturer of the dosage form.
29 January 1996	Change of specification of the active substance.