Post Authorisation Assessments

Chlorsol 50% Powder for Oral Solution
Vm 08007/4052

•	17 May 2019	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
•	02 May 2018	Change in the address of the marketing authorisation holder from Vetoquinol UK Limited, Vetoquinol House, Great Slade, Buckingham Industrial Park, Buckingham, MK18 1PA to Vetoquinol UK Limited, Steadings Barn, Pury Hill Business park, Nr. Alderton, Towcester, Northamptonshire, NN12 7LS.
•	15 June 2016	Submission of an updated certificate of suitability.
•	24 May 2016	Change in batch size of the finished product.
•	27 December 2012	Change of packaging not in contact with the finished product formulation
•	29 November 2012	Submission of a new Ph. Eur. Certificate of Suitability for an active substance from a previously unapproved manufacturer Submission of an updated Ph. Eur. Certificate of Suitability for an active substance from a previously approved manufacturer Change of specification of a starting material used in the manufacture of the active substance
•	15 December 2009	Updated of part II of the dossier
•	07 May 2009	Addition of a site for batch release
•	25 October 2006	Update of SPC and Product Literature to bring in line with new legislation
•	31 March 2006	Renewal
•	22 June 2005	Harmonisation of SPC Change of MAH address
•	10 June 2005	Change of manufacturing site including batch release
•	25 March 2004	Change of site of final product testing
•	12 December 2003	Addition of a manufacturer of the dosage form
•	21 November 2003	Addition of a manufacturer of the dosage form
•	17 October 2003	Renewal
•	02 May 2003	Change of manufacturer of the active substance
•	06 August 1997	Change of MAH
•	25 March 1996	Change in size of non-sterile containers