



Post Authorisation Assessments

Anivit B12 Solution for Injection 1000µg/ml

•	26 February 2019	Submission of an updated Ph. Eur. certificate of suitability from an already approved manufacturer.
•	12 February 2019	Changes to an existing pharmacovigilance system as described in the DDPS.
•	03 November 2016	Change in the address of the Marketing Authorisation Holder.
•	07 May 2015	Submission of an updated certificate of suitability.
•	25 February 2013	Change in supplier of packaging component.
•	14 February 2013	Submission of a new Ph. Eur. Certificate of Suitability for an active substance.
•	09 February 2010	Change of MAH from Dales Pharmaceuticals to Dechra Limited.
•	15 January 2009	Renewal.
•	15 August 2007	Change in legal category from PML to POM-VPS Changes to the SPC and Product Literature to bring in line with new legislation.
•	15 August 2005	Renewal.
•	11 November 2004	Change to bring SPC and product literature in line with new legislation.