



Post Authorisation Assessments

Quiflor Multi 100 mg/ml Solution for Injection for Cattle and Pigs (Sows) Vm 01656/4055

•	02 March 2022	Submission of a new Ph. Eur. certificate of suitability for an active substance.
•	15 February 2022	Minor changes to an approved test procedure of the finished product.
•	20 December 2018	Change of distributor details, from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to Alloga UK Limited, Centaur Services Limited, National veterinary Services Limited, Henry Schein UK Holdings Limited. Changes to the labelling and package leaflet. Change in local representative.
•	20 December 2018	Change in the invented name of the veterinary medicinal product from Ubiflox to Quiflor Multi.
•	02 August 2016	Extension of retest period of active substance.
•	09 June 2016	Renewal – UK as RMS
•	20 May 2015	Change in manufacturing site of the active substance.
•	25 November 2013	Change of distributor.
•	05 April 2013	Extension of shelf life of finished product from 2 to 3 years.
•	08 February 2013	Addition of a third route of synthesis (ROS3) for the active substance and deletion of a manufacturing site for ROS1.
•	06 December 2012	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS .
•	20 September 2012	Addition of a new withdrawal period. To add an additional indication. To add an additional dosage regimen.
•	25 July 2012	To change the MAH (from Milklich Laboratorios to Krka, d.d) and Vm numbers.