

Post Authorisation Assessments

Ubrolexin Intramammary Suspension for Lactating Dairy Cows Vm 08327/5009

•	06 April 2023	Change in the name or address or contact details						
		a qualified person for pharmacovigilance.						
•	28 September 2022	Substantial updates to an ASMF.						
•	28 September 2021	Deletion of a non-significant parameter of an active						
		substance.						
•	05 February 2021	Submission of an updated Ph. Eur. certificate of						
		suitability for an active substance from an already						
		approved manufacturer.						
•	24 September 2019	Change in the safety database of an existing						
		pharmacovigilance system as described in the						
		DDPS.						
•	23 May 2019	Extension of a storage period of the active						
		substance.						
•	06 March 2019	Change in the QPPV of an existing						
		pharmacovigilance system as described in the						
	07.01	DDPS.						
•	07 November 2018	Change of MAH, from Boehringer Ingelheim Ltd,						
		Ellesfield Avenue, Bracknell, Berkshire, RG12 8YS						
		to Boehringer Ingelheim Animal Health UK Ltd,						
	00.4	Ellesfield Avenue, Bracknell, Berkshire, RG12 8YS.						
•	30 August 2017	Deletion of manufacturing site for an active						
		substance.						
		Deletion of manufacturing site for an active						
		substance.						
	22 December 2015	Addition of a manufacturer of the active substance.						
•	23 December 2015	Changes to DDPS following the assessment of the						
		same DDPS in relation to another medicinal product of the same MAH						
	24 April 2013	Change in name of the manufacturer of the finished						
•		product.						
•	24 September 2012	Renewal procedure – Ireland as RMS.						
	11 January 2012							
•	11 January 2012	To change the address of the MAH in Spain and Portugal.						
•	11 January 2012	To change the address of the MAH in Italy.						
•	23 September 2010	Addition of a manufacturing site and a milling site for						
		the active substance. Minor change to						
		manufacturing process of active substance.						
•	23 September 2010	Change in the re-test period/storage period or						
		storage conditions of the active substance.						
•	23 September 2010	Change to in-process tests or limits applied during						
		the manufacture of the finished product.						
L	1							

•	3 September 2010	Change	in	the	batch	size	(including	batch	size		
	ranges) of the finished product.										