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## Post Authorisation Assessments

•	27 September 2018	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
		Change in the contact details of the QPPV of an existing
		pharmacovigilance system as described in the DDPS.
		Change of the back-up procedure of the QPPV of an
		existing pharmacovigilance system as described in the DDPS.
•	26 February 2015	Change in distributor details.
•	21 June 2013	Change to the MAH address.
•	19 September 2008	Changes to bring SPC and Product Literature in line with
		new legislation and change of legal category from POM to POM-V.
•	14 December 2005	Deletion of indications.
•	14 December 2005	Renewal.
•	05 November 2004	Changes to bring SPC and Product Literature in line with new legislation.
•	26 February 2004	Renewal.
•	28 November 2001	Change to manufacturing site of the dosage form.
•	11 October 2001	Change to specification of the finished product.
•	13 September 2001	Change to name of manufacturing site.
•	16 February 2001	Change to manufacturer of the active substance.
•	07 May 1999	Change in name of product from 'Atrocare' to 'Atrocare Injection'.
•	19 January 1998	Change to the name of MAH from 'Animal Care Ltd' to 'Animalcare Ltd' and change in name of product from 'Atrocare 600µg Solution for Injection' to 'Atrocare'.
•	20 November 1995	Change to dosage and administration details.
•	02 May 1995	Name of product sold as alternate presentation.

## Atrocare 600µg/ml Solution for Injection