Post Authorisation Assessments

Oxycare Tablets 100 mg

•	26 February 2015	Change in distributor details.
•	04 June 2013	Change in the address of the MAH
•	14 February 2012	Submission of an updated EDQM certificate of suitability for the active substance.
•	13 May 2009	Change in the manufacturer of the active substance.
•	19 December 2008	Change in the name/address of a manufacturer of the finished product.
•	30 September 2008	Changes to the SPC and product literature to bring them into line with new legislation.
•	30 September 2008	Change in the legal category from POM to POM-V.
•	21 May 2007	Renewal
•	25 February 2003	Renewal
•	19 April 2002	Deletion of a manufacturer of the active substance.
•	31 July 2001	Increase in tablet weight.
•	31 July 2001	Addition of a manufacturer and assembler of the finished product.
•	28 June 2001	Change in manufacturing process of the active substance.
•	07 November 2000	Change to colourant.
•	07 November 2000	Change to finished product specification.
•	06 December 1999	Addition of a site of assembly of the finished product.
•	18 February 1998	Change to finished product specification.
•	24 September 1997	Change in the name of the product.
•	22 June 1997	Renewal