Post Authorisation Assessments

Oxycare Tablets 50 mg

•	26 February 2015	Change in distributor details.
•	04 June 2013	Change to the address of the MAH
•	14 February 2012	Submission of an updated EDQM certificate of suitability of the active substance.
•	13 May 2009	Change in the manufacturer of the active substance.
•	19 December 2008	Change in the name/address of the finished product manufacturer.
•	17 September 2008	Changes to the SPC and product literature to bring them into line with new legislation.
•	17 September 2008	Change in the legal category from POM to POM-V.
•	01 November 2006	Renewal
•	23 January 2004	Renewal
•	19 April 2002	Deletion of an active substance manufacturer.
•	28 September 2001	Change in tablet weight to meet British Pharmacopoeia requirements.
•	30 July 2001	Addition of a manufacturer and assembler of the finished product.
•	21 June 2001	Change to the reference DMF.
•	30 October 2000	Change in formulation.
•	30 October 2000	Change in finished product specification.
•	06 December 1999	Change in assembler of the finshed product.
•	03 March 1998	Renewal
•	18 February 1998	Change to finished product specification.
•	24 September 1997	Change in the name of the finished product.
•	22 June 1997	Renewal
•	03 March 1995	Change if the manufacturer of the finished product.