



Post Authorisation Assessments

Vetoryl 120 mg Hard Capsules Vm 10434/3002

•	19 March 2024	Change in the address or contact details of a manufacturer or supplier of the starting material, used in the manufacture of the active substance where no European Pharmacopoeia (Ph. Eur.) Certificate of Suitability (CEP) is part of the approved dossier. Change to comply with Ph. Eur. for the starting material used in the manufacture of the active substance. Changes to the quality part of the dossier: Deletion of a test procedure for a reagent used in the manufacture of the active substance. Changes to the quality part of the dossier: Deletion of - a non-significant specification parameter of an active substance.
•	27 February 2024	Change in qualitative composition of the immediate packaging for a solid pharmaceutical form for a finished product.
•	28 November 2023	One-off alignment of the product information with version 9.0* of the QRD template.
•	16 November 2023	Minor changes – to an approved test procedure for active substance, for the finished product, for the immediate packaging of the active substance or the finished product, or of a measuring or administration device.
•	07 September 2022	Deletion of a non-significant in-process test during the manufacture of the finished product.
•	28 July 2022	Editorial changes to Part II B of the dossier for the 10 mg hard capsules. Editorial changes to Part II C of the dossier for the 5 mg and 60 mg hard capsules. Editorial changes to Part II F of the dossier for the 30 mg, 60 mg and 120 mg hard capsules.
•	29 March 2022	Deletion of a non-significant in-process test applied during the manufacture of the finished product.
•	16 November 2021	Addition of a new specification parameter to the specification with its corresponding test method of the finished product.
•	23 September 2021	Addition of a primary packaging site of the finished product. Addition of a secondary packaging site of the finished product.
•	09 February 2021	Submission of an updated Ph. Eur. TSE certificate of suitability for a starting material from an already approved manufacturer.
•	21 April 2020	Deletion of manufacturing site where batch control takes place.
•	17 October 2019	Submission of an updated Ph. Eur. TSE certificate of suitability for a starting material from an already approved manufacturer. Submission of a new Ph. Eur. TSE certificate of suitability from a new / already approved manufacturer. Deletion of Ph. Eur. TSE certificates of suitability.

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•	24 September 2019	Changes to a test procedure for the finished product.
•	07 August 2019	Update to the ASMF.
•	18 June 2019	Addition of a manufacturer responsible for batch release including batch control/testing.
•	02 April 2019	Minor change to an approved test procedure for the active substance.
•	12 February 2019	Changes to an existing pharmacovigilance system as described in the DDPS.
•	08 February 2019	Introduction of a new site of manufacture.
•	01 August 2018	Change in RMS from UK to IE.
•	02 February 2017	Changes to the labelling and package leaflet.
•	17 May 2016	Deletion of a TSE certificate. Submission of a new TSE certificate. Submission of an updated TSE certificate. Submission of a new TSE certificate. Submission of a new TSE certificate.
•	25 November 2015	Additional site for batch testing of the finished product.
•	21 May 2015	Submission of updated Ph. Eur. Certificates of Suitability from an already approved manufacturer
•	01 May 2015	Change in name of a manufacturer of the active substance
•	20 November 2014	Updates to the labelling and package leaflet.
•	10 October 2014	Change of MA holder address.
•	06 March 2014	Significant change to the SPC with regard to clinical data.
•	14 November 2013	Grouped variation to update the DMF, amend the raw materials specifications, update the manufacturing methods, and to optimise the active substance manufacturing method.
•	03 May 2012	Grouped variation to update TSE Certificates of Suitability as well as the addition of a new Certificate of Suitability for an active substance manufacturer.
•	02 March 2012	Change in the manufacturer of the active substance.
•	09 December 2011	To change the ink used to mark the capsules.
•	01 April 2011	Changes to an existing pharmacovigilance system as described in the DDPS.
•	15 December 2010	Change of Distributor.
•	06 October 2010	To change the Marketing Authorisation Holder.
•	17 September 2010	Change in immediate packaging of the finished product.
•	17 September 2010	Submission of new or updated Ph. Eur Certificates of Suitability.
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•	07 June 2010	Renewal
•	03 October 2008	Change active/intermediate batch size.
•	05 July 2007	Change test procedure for active/active component .
•	01 June 2007	Change shelf life of finished product as packaged for sale.
•	01 June 2007	Change finished product test procedure.

•	01 June 2007	Change finished product test procedure.
•	16 May 2007	Change/addition of imprints/bossing/markings on tabs or capsules.
•	08 March 2006	Change of distributor.