Veterinary Medicines Directorate

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Post Authorisation Assessments

Senvelgo 15 mg/ml Oral Solution for Cats

Vm 08327/5004

05 August 2026	Addition to a quality control testing site for the finished product. Addition to a quality control testing site for the finished product
25 June 2026	Indication changed to: 'For the reduction of hyperglycaemia in cats with non-insulin-dependent diabetes mellitus', addition of vomiting as an AE, changes to sections 3.5 and 3.6.
19 March 2026	Minor changes in the manufacturing process of an active substance.
15 January 2026	Addition of a manufacturer responsible for batch control arrangements and quality testing of the finished product. Addition of a new in-process test and limits applied during the manufacture of the finished product. Minor changes to production equipment when described in the dossier. Addition of a manufacturer responsible for batch release including batch control or testing of a sterile or non-sterile finished product. Addition of a primary packaging site of a non-sterile finished product. Addition of a secondary packaging site of a finished product.
15 January 2026	Addition of a manufacturing site for part or all of the manufacturing process of the finished product. Changes in the composition (excipients) of the finished product. Minor change in the manufacturing process of the finished product.
17 September 2025	Change in test procedure for a starting material used in the manufacturing process of the active substance.
31 July 2025	Minor changes: to an approved test procedure for active substance.
26 January 2025	Additional pack size: 35 mL bottle and a nominal fill volume of 12 mL. Alignment of the product information with version 9.0* of the QRD templates.
07 October 2024	Deletion of a manufacturing site where batch control takes place.
10 July 2024	Minor changes in the manufacturing process of an active substance.
06 July 2024	Addition of a manufacturer for an active substance intermediate.
13 April 2024	Minor changes to an approved test procedure for a starting material used in the manufacturing process of the active substance.
16 January 2024	Change in the re-test period of the active substance where no

	Ph. Eur. Certificate of Suitability covering the retest period is part of the approved dossier.