



Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021,
please refer to the [EMA](#) website.

Zuprevo 180 mg/ml Solution for Injection for Cattle

Vm 01708/5060

09 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
22 February 2026	Change in the manufacturer of the active substance: - Introduction of a manufacturer of the active substance supported by an ASMF.
01 December 2024	Minor change to an approved test procedure for the finished product.
08 January 2024	Change(s) in the SPC, labelling or package leaflet intended to implement the outcome of a procedure or recommendations from the competent authority or the Agency concerning risk management measures in pharmacovigilance related to veterinary medicinal products. One-off alignment of the product information with version 9.0* of the QRD templates
02 November 2022	Minor changes in the restricted part of ASMF. Minor changes in the restricted part of ASMF. Minor changes in the applicants part of ASMF.
03 March 2022	Minor change in the manufacturing process.
24 February 2022	Changes to the labelling and/or package leaflet.
17 December 2021	Addition of a new specification parameter with its corresponding test method of an intermediate used in the manufacturing process of the active substance. Minor change to an approved test procedure used in the manufacturing process of the active substance. Addition of a new in-process test and limit applied during the manufacture of the active substance. Changes to a test procedure for the starting material.
23 August 2021	Extension of re-test period of the active substance.