



Post Authorisation Assessments

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

Eurican Herpes 205 Powder and Solvent for Emulsion for Injection Vm 04491/5008

•	31 October 2023	Update to the description of starting materials of biological origin.
•	10 October 2023	Minor changes to the production equipment.
•	11 April 2023	Change in the name or address or contact details of a qualified person for pharmacovigilance.
•	20 December 2022	To add use of recombinant trypsin as a substitute to porcine trypsin for the manufacture of the active substance.
•	18 October 2022	Lower osmolality specification of stabiliser 49 used as a freeze-drying substrate changed to 320 mOsm/kg.
•	15 December 2021	Changes to the labelling and/or package leaflet.
	25 March 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	19 March 2021	Minor changes to an approved test procedure of the finished product