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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

•	20 November 2024	Alignment of the SPC/QRD text with the newest EU version 9.0 QRD template and GB National SPC/QRD template.
•	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