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# Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021, please refer to the EMA website.

## Purevax RC Lyophilisate and Solvent for Suspension for Injection

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| • | 03 November 2022 | Editorial change of the range for osmolality for stabiliser 49 to 320–356 mOsm/kg. |
| • | 18 August 2022 | Deletion of a manufacturer of an active substance. |
| • | 16 August 2022 | Modification of the conductivity test limit of acceptance carried out on sterile diluent to less than or equal to 25 µS/cm. |
| • | 05 August 2022 | To increase the maximum release titre of the feline herpesvirus component of Purevax vaccines from 10^6.2 CCID50/dose to 10^6.5 CCID50/dose and to align the titre across the documents to be expressed in units/dose. |
| • | 06 May 2022 | Changes to SPC & product literature following a periodic safety update report (PSUR) |
| • | 17 March 2022 | Change of a test procedure for the active substance.<br>Changes in the manufacturing process of the active substance. |
| • | 25 June 2021 | Addition of a new specification parameter to the specification with its corresponding test method of the finished product.<br>Deletion of a non-significant specification parameter of the finished product. |
| • | 26 March 2021 | Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. |