21 February 2020
EMA/CVMP/624348/2019
Committee for Medicinal Products for Veterinary Use

Summary of opinion[1] (post-authorisation)

Clynav

Common name: pUK-SPDV-poly2#1 DNA plasmid coding for salmon pancreas disease virus proteins

On 20 February 2020, the Committee for Medicinal Products for Veterinary Use (CVMP) adopted a positive opinion[2], recommending the granting of a variation to the terms of the marketing authorisation for the veterinary medicinal product Clynav. The marketing authorisation holder for this veterinary medicinal product is Elanco GmbH.

Clynav is currently authorised as a solution for injection. The variation concerns extending the duration of immunity from 3 months after vaccination to 12 months after vaccination.

Detailed conditions for the use of this product are described in the updated summary of product characteristics (SPC), for which an updated version reflecting the changes will be published in the revised European public assessment report (EPAR) and will be available in all official European Union languages after the variation to the marketing authorisation has been granted by the European Commission.

[1] Summaries of opinion are published without prejudice to the Commission Decision.
[2] Applicants may appeal any CVMP opinion, provided they notify the European Medicines Agency in writing of their intention to appeal within 15 days of receipt of the opinion.

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union