



Post Authorisation Assessments

Gallifen 40 mg/g Premix for Medicated Feeding Stuff for Chickens and Pheasants

Vm 30282/4029

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| • | 05 September 2023 | Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance, – starting material, reagent or intermediate used in the manufacturing process of the active substance, or – excipient. (GB) |
| • | 16 December 2021 | Renewal – UK as CMS. |
| • | 13 August 2021 | Submission of a new Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. |
| • | 22 November 2019 | Increase in the shelf-life of the finished product after dilution, from 1 month to 3 months. |
| • | 30 May 2019 | Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer. |
| • | 22 February 2019 | Change in storage conditions and shelf life of the finished product. |
| • | 15 November 2018 | Addition of pheasants as a new target species. |
| • | 06 September 2018 | Change in the SPC, labelling or package leaflet due to new data. |