



Post Authorisation Assessments

Dermipred 20 mg Tablets for Dogs Vm 15052/5079

• 09 December 2024	Reduction in the shelf-life from 3 years to 2 years for the finished products packaged in Al/PVC –Al-OPA blisters and change in storage conditions reducing maximum storage temperature from 30C to 25C.
• 16 August 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance.
• 22 December 2022	Change in the re-test period/storage period of the active substance. Submission of a new Ph. Eur. certificate of suitability for an already approved active substance manufacturer.
• 30 September 2022	Change of MAH address from: Unit 3 Anglo Office Park, White Lion Road, Amersham, Buckinghamshire, HP7 9FB to Explorer House, Mercury Park, Wycombe Lane, Wooburn Green, High Wycombe, Buckinghamshire, HP10 0HH, United Kingdom.
• 08 October 2021	Renewal – UK as CMS.
• 24 December 2019	Change in the invented name of the veterinary medicinal product from Prednisolone Ceva 10 mg tablets for dogs to Dermipred 10 mg tablets for dogs in France.
• 04 November 2019	Increase in the shelf-life of the finished product as packaged for sale, from 2 years to 3 years.
• 23 May 2019	Replacement of a site where batch control/testing takes place.
• 21 February 2018	Repeat Use application to add 3 new member states.
• 19 September 2017	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.