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Post Authorisation Assessments

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

Cerenia 10 mg/ml Solution for Injection for Dogs and Cats Vm 42058/5008

• Ma	ay 2024	Deletion of a non-significant specification parameter in the specification parameters of an excipient.
• 28	April 2024	Deletion of a manufacturing and primary packaging site of the finished product.
• 20	February 2024	One-off alignment of the product information with the latest QRD template.
• 19	January 2024	Minor change in the manufacturing process of the active substance.
• 28	April 2023	Deletion of a manufacturer of the finished product.
• 31	October 2022	Change in name and address of a manufacturer of the active substance.
• 02	September 2022	Change(s) in the SPC, labelling or package to section 4.6 and 6.
• 30	December 2021	To implement changes to the SPC and package leaflet following assessment of the PSUR.
• 16	December 2021	Change to part of the (primary) packaging material not in contact with the finished product formulation. Change to part of the (primary) packaging material not in contact with the finished product formulation.
• 08	April 2021	Minor change to an approved test procedure for the active substance used in the manufacturing process of the active substance.