Veterinary Medicines Directorate

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Post Authorisation Assessments

Cazitel Plus Tablets for Dogs

Vm 08749/3002

19 January 2025	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance (NI) Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance (NI) Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance (NI)
10 April 2024	Deletion of a manufacturer of the active substance.
22 August 2023	SRP to add one member state.
20 June 2023	One-off alignment of the product information with version 9.0* of the QRD templates i.e. major update of the QRD templates in accordance with Regulation (EU) 2019/6, for veterinary medicinal products placed on the market in accordance with Directive 2001/82/EC or Regulation (EC) No 726/2004.
08 June 2023	Addition of a manufacturer of the active substance.
12 May 2023	Substantial changes in the updated version of the ASMF or the active substance part of the dossier.
23 January 2023	Change in product name from Cazitel Plus XL, Comprimate to Prazitel Plus XL, Comprimate in RO only.
20 April 2022	Update to ASMF.
06 May 2021	Change in distributor from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP.
22 April 2021	Deletion of manufacturing site for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
08 October 2019	Increase in the shelf-life of the finished product as packaged for sale, from 3 years to 5 years.
15 July 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
10 January 2019	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
02 July 2018	ASMF updated.
22 December 2017	Addition of a new Ph. Eur. certificate of suitability for an

	active substance from a new manufacturer.
15 May 2015	Submission of a new certificate of suitability.
01 October 2014	Change to the product name in Sweden only, from ‘Cazitel, 150mg/144mg/50mg tablet för hund’ to ‘Cazitel comp 150 mg/144 mg/50 mg tablett för hund’
21 July 2014	Change of distributor from Pfizer Ltd. to Zoetis UK Limited.
18 July 2014	Renewal procedure – Ireland as RMS.
10 June 2013	To add an additional active substance manufacturer.
01 February 2013	Submission of an updated Ph. Eur. Certificate of Suitability for an already approved manufacturer.
02 May 2012	Submission of a new or updated Ph. Eur. Certificate of Suitability.
26 October 2011	Repeat Use Comm.
02 July 2010	To add a pork flavour to the tablets.
12 May 2010	To change the colour of the mock-ups from blue to yellow.