Veterinary Medicines Directorate

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Post Authorisation Assessments

Clavucill 400 mg/100 mg Tablets for Dogs

Vm 19968/4004

15 October 2026	Other variations not specifically covered elsewhere in chapter G which involve the submission of studies to the competent authority, including additional clinical and non-clinical studies, including BE-studies. One-off alignment of the product information with version 3 of the GB QRD templates.
18 February 2026	Submission of a Ph. Eur. CEP for an active substance. Submission of a Ph. Eur. CEP for an active substance. Deletion of a manufacturer of an active substance.
19 November 2024	Change in the address of a manufacturer of the finished product.
15 July 2022	Updated certificate of suitability for an already approved manufacture. Updated certificate of suitability for an already approved manufacture. New certificate of suitability from a new manufacturer for an active substance. New certificate of suitability from a new manufacturer for an active substance.
27 December 2018	Changes to the labelling and package leaflet.
13 November 2018	Change in distributor details from: Chenelle Animal Heath Ltd., 7 Rodney Street, Liverpool, L1 9HZ, UK, to V.M.D. n.v., Hoge Mauw 900, 2370 Arendonk, Belgium.
02 October 2018	Change in shape or dimensions of the container or closure (immediate packaging).
17 November 2016	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
30 July 2015	Introduction of a new pharmacovigilance system, which has not been assessed by the relevant NCA/EMA.
03 June 2015	Change in the name of the medicinal product from Clavucill Tablets 500 mg to Clavucill 400 mg/100 mg, Tablets for dogs.
25 March 2015	Change to the MAH.
12 February 2014	Mock-ups updated in line with Renewal procedure. Changes not connected with SPC.
28 January 2014	Submission of new or updated Certificates of Suitability.
05 July 2013	Renewal procedure.
19 May 2009	To update the reference number of the certificate of suitability for Potassium Clavulanate.