

Post Authorisation Assessments

Clavucill 400 mg/100 mg, Tablets for Dogs Vm 19968/4004

• 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.
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		manufacturer for an active substance.
• 27 Dece	ember 2018	Changes to the labelling and package leaflet.
• 13 Nove	ember 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 Octo	ber 2018	Change in shape or dimensions of the container
		or closure (immediate packaging).
• 17 Nove	ember 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	e 2015	Change in the name of the medicinal product
		from Clavucill Tablets 500 mg to Clavucill 400
05.14	1 0045	mg/100 mg, Tablets for dogs.
• 25 Marc		Change to the MAH.
• 12 Febr	uary 2014	Mock-ups updated in line with Renewal
	0044	procedure. Changes not connected with SPC.
• 28 Janu	ary 2014	Submission of new or updated Certificates of
	0040	Suitability.
• 05 July		Renewal procedure.
• 19 May	2009	To update the reference number of the certificate
		of suitability for Potassium Clavulanate.