Veterinary Medicines Directorate

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Post Authorisation Assessments

Canergy 100 mg Tablets for Dogs

Vm 41821/4019

•	04 May 2024	Addition of an alternative artificial beef flavour in the manufacture of the finished product. (GB)
•	04 May 2024	Addition of use of the non-pharmacopoeial excipient artificial beef flavour synthesized without materials of animal origin and complying to EU food guideline EC/1334/2008. (NI only - Refused for GB) Add an alternative site responsible for batch release of the finished product. (NI+GB) Add an alternative site responsible for primary packaging of the finished product. (NI+GB) Add an alternative site responsible for secondary packaging of the finished product. (NI+GB)
•	04 May 2024	To add medium density polyethylene 60 um bags for storage of the bulk tablets. (NI+GB) To include a hold time for the bulk tablets as stored in MDPE 60 µm bags. (NI+GB) To change the bulk and tapped density in-process limits applied during the manufacture of the finished product. (NI+GB) Add an alternative site responsible for manufacturing of the finished product. (NI+GB)
•	26 May 2021	Repeat Use application to add 1 new member state.
•	18 May 2020	Renewal - UK as CMS.
•	15 August 2019	Introduction of a new pharmacovigilance system.
•	21 March 2019	Change of RMS from UK to NL
•	21 December 2015	Change of UK distributor Approval of joint-labelled mock-ups
•	13 July 2015	Additional manufacturing site for secondary packaging and batch release.