Veterinary Medicines Directorate

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Post Authorisation Assessments

Aloquan, 18.7 mg/g + 140.3 mg/g Oral Paste for Horses

•	27 October 2020	Addition of a secondary packaging site of the finished product.
•	20 December 2019	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.
•	11 September 2019	Addition of a manufacturer responsible for batch release of the finished product.
•	30 July 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	18 October 2018	Renewal - UK as CMS
•	07 February 2018	Deletion of manufacturing site for an active substance.
•	11 April 2017	Submission of a new certificate of suitability. Submission of a new certificate of suitability.
•	23 December 2015	Change of Distributor from Farm & Stable Supplies LLP to GBA International
•	31 July 2015	Submission of an updated Ph. Eur. Certificate of Suitability.
•	22 July 2015	Change to the product name in Sweden only.
•	29 August 2014	Changes to the DDPS.
•	12 June 2014	Update to the European Pharmacopoeial Certificate of Suitability.
•	08 May 2014	To change the product's trade name in the United Kingdom (to "Aloquan, 18.7mg/g + 140.3mg/g Oral Paste for Horses"), in Germany (to "Aloquan, 18.7mg/g + 140.3mg/g Oral Paste for Horses"), in Denmark (to "Noropraz Vet, 18.7mg/g + 140.3mg/g Oral Paste for Horses"), and in Sweden (to "Ivermectin/Praziquantel Norbrook,18.7mg/g + 140.3mg/g Oral Paste for Horses").
•	24 April 2014	Change to distributor details, from Norbrook Laboratories (GB) Limited to Farm & Stable Supplies LLP.