



Post Authorisation Assessments

Johnson's Easy Spot-On Solution Wormer for Cats & Kittens 20mg Vm 08007/4171

27 February 2025	Update of mock ups.
09 October 2024	Slight rewording of section 4.6.
09 October 2024	Downscaling of finished product batch size. Addition of new in-process tests applied during finished product manufacture. Deletion of a non-significant in-process test during the manufacture of the finished product. Minor change to an approved test procedure for the active substance.
09 October 2024	Change to test procedure for the immediate packaging of the finished product. Addition of a new specification parameter for the finished product immediate packaging. Addition of a batch control and quality testing site for the finished product. Deletion of obsolete specification parameters for the finished product immediate packaging. Deletion of obsolete specification parameters for the finished product immediate packaging. Deletion of obsolete specification parameters for the finished product immediate packaging. Deletion of Ph. Eur. CEP for an active substance. Addition of a manufacturer responsive for batch release of the finished product. Addition of a primary packaging site for the finished product. Addition of a secondary packaging site for a finished product.
09 October 2024	Addition of a test procedure for the finished product. Change in the specification parameters of the finished product. Change to in-process limit applied during the manufacture of the finished product. Addition of a manufacturing site for the finished product.
09 October 2024	To introduce information on the packaging regarding the risks posed by the excipient N-methyl pyrrolidone.
20 October 2022	Updated certificate of suitability from an already approved manufacturer.
19 October 2021	Introduction of a new pharmacovigilance system.
05 November 2020	Change of MAH from Bayer Animal Health GmbH, 51368 Leverkusen, Germany to Vetoquinol UK Limited, Steadings Barn, Pury Hill Business Park, Nr. Alderton,

	Towcester, Northamptonshire, NN12 7LS.
09 June 2020	Change of MAH, from Bayer plc, 400 South Oak Way, Green Park, Reading, Berkshire, RG2 6AD to Bayer Animal Health GmbH, 51368 Leverkusen, Germany.
09 June 2020	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.
03 March 2020	Submission of a new Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
17 September 2019	Change in the invented name of the veterinary medicinal product from Bob Martin Spot On Solution Dewormer 20 mg to Johnson's Easy Spot-On Solution Wormer for Cats and Kittens 20mg. Changes to the labelling* and/or package leaflet*
16 May 2019	Change in distributor details from Bob Martin (UK) Ltd to Johnson's Veterinary Products Ltd.
18 September 2018	Submission of a new Ph. Eur. certificate of suitability for an active substance excipient from a new manufacturer.
18 September 2018	Change in distributor details from Bayer plc, Animal Health Division, Strawberry Hill, Newbury, Berkshire, RG14 1JA to Bayer plc, 400 South Oak Way, Green park, Reading, Berkshire, RG2 6AD.
05 May 2017	Change in the address of the marketing authorisation holder from Bayer plc, Animal Health Division, Bayer House, Strawberry Hill, Newbury, Berkshire, RG14 1JA to Bayer plc, 400 South Oak Way, Green Park, Reading, Berkshire, RG2 6AD.
27 May 2016	Delete Unidrug Distribution Group Limited as a distributor.
8 December 2015	Addition of a secondary packaging site.
05 February 2015	Reduction in shelf-life of the finished product, from 5 years to 30 months. Addition of specification parameters for the finished product.
31 July 2013	Deletion of 2 non-significant parameters of an active substance used in the manufacturing process of the active substance.
07 September 2011	Update of part of the dossier
16 February 2011	Change of distributor to Unidrug Distribution Group Ltd. And Bob Martin UK Ltd.
02 July 2009	Renewal
23 June 2009	Submission of 2 new/updated Ph. Eur. Certificates of Suitability for an active substance or starting material /reagent/intermediate in the manufacturing process of the active substance.
29 August 2008	Variation to bring the SPC and labels in line with the new legislation and to transfer the legal category from GSL to AVM-GSL.
14 July 2005	Corrections/simple layout changes to package leaflet and carton.

