



Post Authorisation Assessments

Johnson's Wormer Plus 230 mg/20 mg Film-coated Tablets for Cats and Kittens Vm 08007/5049

03 February 2026	Submission of a Ph. Eur. CEP for an active substance.
07 October 2025	Change in pack size (number of tablets in a pack) within the range of the currently approved pack size.
07 October 2025	Change in pack size of the finished product. Change in legal (distribution) category of the VMP. One-off alignment of the product information with version 9.0* of the QRD templates.
13 January 2025	Change in pack size within the range of the currently approved pack size. Change in the name of the veterinary medicinal product from Johnson's One Dose Easy Wormer 230 mg/20 mg Film-coated Tablets for Cats and Kittens to Johnson's Wormer Plus 230/20mg Film-coated Tablets for Cats & Kittens.
17 October 2024	Deletion of one of the authorised immediate packaging forms of the finished product that does not delete a strength or pharmaceutical form. Addition of a manufacturer responsible for batch release including batch control or testing of the finished product. Addition of a primary packaging site of the finished product. Addition of a secondary packaging site of the finished product.
17 October 2024	Change in coating weight of oral dosage forms or change in weight of capsule shells for a solid oral pharmaceutical form. Change in test procedure for the immediate packaging of the finished product. Downscaling down to 10-fold compared to the originally approved batch size of an immediate release oral pharmaceutical form. Increase up to 10-fold compared to the originally approved batch size of an immediate release oral pharmaceutical form. Addition of new specification parameters to the immediate packaging specification with the corresponding test method. Change of specifications of an active substance to fully comply with the Ph. Eur. Deletion of Ph. Eur. CEPs for an active substance. Minor changes to an approved test procedure for active substance. Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. Submission of a new Ph. Eur. CEP from a new manufacturer for a non-sterile active substance.
17 October 2024	Minor change in the manufacturing process of the finished

	<p>product.</p> <p>Changes in the composition (excipients) of the finished product.</p> <p>Addition of a manufacturing site for part or all of the manufacturing process of the finished product.</p> <p>Addition of a manufacturing site for part or all of the manufacturing process of the finished product.</p>
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.
14 October 2019	Addition of a Distributor: Johnson's Veterinary Products Limited, 5 Reddicap Trading Estate Sutton Coldfield, West Midlands, B75 7DF, Tel: 0121 378 1684