



Post Authorisation Assessments

Dycoxon 2.5 mg/ml Oral Suspension for Sheep and Cattle Vm 08749/4085

24 March 2026	Alignment of the product information with version 9.0* of the QRD templates.
15 December 2025	Minor changes - to an approved test procedure for the finished product. (NI) Minor changes - to an approved test procedure for the finished product. (NI)
01 August 2025	Minor changes - to an approved test procedure for the finished product. (GB) Minor changes - to an approved test procedure for the finished product. (GB)
18 January 2025	Deletion of a site where batch control takes place. (GB and NI).
31 August 2022	Unlimited renewal.
28 May 2020	Changes to the labelling and/or package leaflet.
11 April 2019	Change in the invented name of the veterinary medicinal product from Dycoxon 2.5 mg/ml Oral suspension for Sheep and Cattle to Diacox 2.5 mg/ml Oral suspension for Sheep and Cattle in FR and DE only.
26 March 2019	Change of RMS from UK to ES.
15 March 2019	Updated of the ASMF.
12 February 2019	Change in the name of the medicinal product in Spain, Portugal and Italy and an editorial change to the QRD text for the immediate packaging (inclusion of indications) as requested by another Member State.