



Post Authorisation Assessments

Linco-Sol 400 mg/g Powder for Use in Drinking Water for Pigs and Chickens Vm 32823/4005

•	08 September 2022	New certificate of suitability from a new manufacturer.
•	02 July 2021	Minor changes to an approved test procedure of the finished product. Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State. Change in the SPC, labelling or package leaflet following assessment of the same change for the reference product.
•	26 March 2020	Change in the address of the marketing authorisation holder from Lavet Pharmaceuticals Ltd., Ottó u. 14., Budapest 1161. to Lavet Pharmaceuticals Ltd., 2143 Kistarcsa, Batthyany u. 6., Hungary. Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
•	13 December 2019	Deletion of a manufacturing site for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Introduction of a re-test period of the active substance.
•	05 July 2018	Repeat Use application to add 1 new member state.
•	08 June 2018	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change in the SPC, labelling and package leaflet following assessment of the same change for the reference product.
•	August 2016	Submission of a new certificate of suitability.
•	23 May 2013	Renewal MA
•	15 August 2012	To change the name of the veterinary medicinal product in Poland only, to LincoScan.