



Post Authorisation Assessments

Lotimil Plus 900 mg/33.75 mg Chewable Tablets (> 22–45 kg) Vm 00879/5047

16 February 2026	Deletion of a non-significant specification parameter for an active substance.
16 February 2026	Deletion of a manufacturing site for the finished product. Deletion of a manufacturing site for the finished product.
09 October 2025	Minor change to an approved test procedure for an active substance.
18 August 2025	Change in the specification parameters for a reagent used in the manufacturing process of an active substance. Change in the specification parameters of a reagent used in the manufacture of an active substance. Change in the specification parameters of a reagent used in the manufacture of an active substance. Minor change to the manufacturing process of an active substance.
04 July 2025	Minor changes to an approved test procedure for the active substance.
04 April 2025	Addition of a manufacturer of a starting material used in the manufacturing process of the active substance.
27 February 2025	Change in the name or address or contact details of a manufacturer or supplier of a starting material. Deletion of a supplier of a starting material.