



Post Authorisation Assessments

Alcide Uddergold Platinum Concentrates (Base and Activator) for Teat Dip Solution for Cattle (Dairy) Vm 04509/5000

29 October 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (NI).
18 February 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI).
27 October 2024	Submission of a new Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance. (GB).
06 June 2024	Replacement of a manufacturer responsible for batch release.
19 January 2024	Deletion of Distributor - Progiene a division of Rumenco, Stretton House, Derby Road, Burton on Trent, Staffordshire, DE13 0DW.
08 June 2023	Change of MAH address from Ecolab Ltd, Lotherton Way, Garforth, Leeds, LS25 2JY, United Kingdom to Ecolab Ltd, Unit 1, Wernddu Court, Caerphilly, CF83 3SG, United Kingdom.
16 November 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
07 November 2019	To amend Distributors from: Ecolab Ltd., Duke Avenue, Stanley Green Trading Estate, Cheadle Hulme, Cheshire, SK8 6RB (UK) and Rosebeck Services, Roseberry Court, Ellerbeck Way, Stokesley, N. Yorkshire, TS9 5QT to: Ecolab Ltd., PO Box 11, Winnington Avenue, Northwich, Cheshire, CW8 4DX and Progiene a division of Rumenco Ltd, Stretton House, Derby Road, Burton on Trent, Staffordshire, DE13 0DW.
28 August 2019	Change of MAH from 'Ecolab Deutschland GmbH' to 'Ecolab Ltd'.
09 January 2019	Change in RMS from UK to IE.
11 October 2017	Deletion of a manufacturing site for the finished product. Deletion of a manufacturing site for an active substance. Deletion of a manufacturing site for an active substance. Deletion of a manufacturing site for an active substance.
28 February 2013	Variation to change the address of the Marketing Authorisation Holder and the address of a manufacturer responsible for batch release.
13 May 2011	Variation to change the name of the veterinary medicinal product in the UK and IE.
15 December 2009	Variation to change the name of the Marketing Authorisation Holder and the manufacturer responsible

	for batch release.
17 September	Renewal (UK as RMS)
08 August 2008	MRP (UK as RMS)
20 July 2007	Change of the active substance manufacturer.
11 August 2006	Deletion of an importer responsible for batch release.
11 August 2006	Deletion of an importer responsible for batch release.
08 August 2006	Variation to change the name of an active substance manufacturer.
04 July 2006	Variation to change the distributor.
16 May 2006	Variation to change Part IV of the dossier.
10 May 2006	Variation to change Part 3B of the dossier.
05 April 2006	Variation to change Part IV of the dossier.
05 April 2006	Variation to change Part IV of the dossier.
27 February 2006	Variation to change the storage conditions of the finished product.
24 January 2006	Variation to update Part III of the dossier.
09 December 2005	Variation to change the name of the veterinary medicinal product.
29 November 2005	Variation to change the name and address of the Marketing Authorisation Holder.
15 November 2005	Variation to include an additional pack size.
15 November 2005	Addition of a manufacturer in the EU.
15 February 2005	Renewal
01 September 2004	Addition of an active substance manufacturer.
27 November 2003	Change in the specification of an excipient.
27 November 2003	Variation to replace an excipient with a comparable excipient.
23 May 2003	Variation to change the formulation of the finished product.
21 February 2003	Change in the qualitative composition of the immediate packaging material.
21 February 2003	Addition of an excipient.
21 February 2003	Change in the finished product excipients (deletion of a colorant).
21 February 2003	Change in the specification of the product.
21 February 2003	Change of an excipient with a comparable excipient.
21 February 2003	Change of name of the veterinary medicinal product.
15 November 2002	Change of manufacturer and assembler of dosage form.
30 August 2002	Change of distributor and importer of the final dosage form from outside EU.
02 August 2001	Change of importer of the final dosage form.
12 January 2001	Renewal.
07 January 1998	Change of manufacturer and assembler of dosage form.
04 June 1997	Change of manufacturer and assembler of dosage form.
05 March 1997	Addition of a finished product manufacturer.
29 January 1997	Change to the ingredient specification.
09 January 1997	Change of address of the active substance manufacturer.
06 December 1996	Addition of an active substance manufacturer.

02 October 1996	Change of address of a manufacturer/assembler.
02 October 1996	Change of Marketing Authorisation Holder.
04 October 1996	Change of address of an importer.
26 July 1996	Change of finished product formulation.
26 July 1996	Change of pack size of the finished product.