Veterinary Medicines Directorate

Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS
Telephone +44 (0)1932 336911

Post Authorisation Assessments

Alcide Uddergold Platinum Concentrates (Base and Activator) for Teat Dip Solution for Cattle (Dairy)

Vm 04509/4014

•	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.