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

Not authorised

- Lactic acid
- Sodium chlorite

Product identification

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

Active substance:
Lactic acid

Sodium chlorite

Target species:
Cattle

Route of administration:
Teat use

Product details

Active substance and strength:
Lactic acid
26.40 milligram(s) / 1.00 millilitre(s)

Sodium chlorite
6.40 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Concentrate for dip solution

Withdrawal period by route of administration:
Teat use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG52A

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:
Surrendered

Authorised in:
United Kingdom (Northern Ireland)

Package description:
Base and Activator are supplied separately, packed in pairs of high density polyethylene (HDPE) bottles/container withpolypropylene or HDPE screw caps holding 20 litres of the Base or Activator component.
Base and Activator are supplied separately, packed in pairs of high density polyethylene (HDPE) bottles/containers withpolypropylene or HDPE screw caps

holding 10 litres of the Base or Activator component.
Base and Activator are supplied separately, packed in pairs of high density polyethylene (HDPE) bottles/containers withpolypropylene or HDPE screw caps holding 3.785 litres of the Base or Activator component.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ecolab Limited

Marketing authorisation date:
25/05/1994

Manufacturing sites for batch release:
Ferdinand Eimermacher GmbH und Co. KG

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 04509/4014

Date of authorisation status change:
6/11/2025

Reference member state:
Ireland

Procedure number:
IE/V/0572/001

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

ie-puar-mr-iev0572001-alcide-uddergold-platinum-en.pdf