Alcide UDDERgold Platinum

Authorised

- Lactic acid
- Sodium chlorite

Product identification

Medicine name:
Alcide UDDERgold Platinum
Alcide UDDERgold Platinum

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

Authorisation status:

Valid

Authorised in:

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 Deutschland GmbH

Marketing authorisation date:
30/04/2008

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10795/003/001

Date of authorisation status change:
30/04/2008

Reference member state:
Ireland

Procedure number:
IE/V/0572/001

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics
