

Alcide UDDERgold Platinum

Not authorised

- Lactic acid
- Sodium chlorite

Product identification

Medicine name:

Alcide UDDERgold Platinum

Alcide UDDERgold PM vet, concentraten (Base en Activator) voor tepeldipoplossing voor rundvee (melkvee)

Active substance:

Lactic acid

Sodium chlorite

Target species:

Cattle

Route of administration:

Topical 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:**Topical 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:

Surrendered

Authorised in:

Netherlands

Package description:

Base and Activator are supplied separately, packed in pairs of high density polyethylene (HDPE) bottles/container with polypropylene 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 with polypropylene 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 with polypropylene 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:

26/05/2009

Manufacturing sites for batch release:

Ecolab Deutschland GmbH

Responsible authority:

MEB

Authorisation number:

REG NL 101795

Date of authorisation status change:

23/03/2023

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

Combined File of all Documents

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