UDDERgold Platinum Germicidal Barrier Teat Dip Solution Concentrates (Base and Activator) for Teat Dip Solution For Cattle (Dairy)

Not authorised

- Lactic acid
- Sodium chlorite

Product identification

Medicine name:
UDDERgold Platinum Germicidal Barrier Teat Dip Solution Concentrates (Base and Activator) for Teat Dip Solution For Cattle (Dairy)

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:
Teat 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

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Authorisation status:
Surrendered

Authorised in:
Ireland

Package description:
Base and Activator are supplied separately, packed in pairs of high density polyethylene (HDPE) bottles/containers 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:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Ecolab Deutschland GmbH

Marketing authorisation date:
11/08/2004

Manufacturing sites for batch release:
Ecolab Deutschland GmbH

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10795/001/001

Date of authorisation status change:
11/08/2004

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