Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml Teat dip/spray solution

Authorised

- Chlorhexidine gluconate

Product identification

Medicine name:
Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml Teat dip/spray solution

Active substance:
Chlorhexidine gluconate

Target species:
Cattle

Route of administration:
Teat use

Product details

Active substance and strength:
Chlorhexidine gluconate
5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Teat dip/spray solution

Withdrawal period by route of administration:

Teat use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD08AC02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Available in:

Bulgaria

Package description:

Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml teat dip/teat spray sol. multidos. cont. 200 l
Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml teat dip/teat spray sol. multidos. cont. 60 l
Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml teat dip/teat spray sol. multidos. cont. 25 l
Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml teat dip/teat spray sol. multidos. cont. 20 l
Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml teat dip/teat spray sol. multidos. cont. 10 l
Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml teat dip/teat spray sol. multidos. cont. 5 l

Kenocidin Spray and Dip Chloorhexidine Digluconaat 5 mg/ml teat dip/teat spray sol. multidos. cont. 1 l

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 8(3) of Directive No 2001/83/EC)

Marketing authorisation holder:
Cid Lines

Marketing authorisation date:
30/06/2015

Manufacturing sites for batch release:
Cid Lines

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2579/01.7.2015

Date of authorisation status change:
30/06/2015

Reference member state:
Belgium

Procedure number:
BE/V/0040/001

Concerned member states:
Bulgaria Cyprus Czechia Estonia France Germany Greece Hungary Ireland Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
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