File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/600000046287

Nafpenzal DC intramamálna suspenzia pre hovädzí dobytok (kravy v období státia na sucho)

Authorised

- Dihydrostreptomycin
- Nafcillin
- Benzylpenicillin procaine

Product identification

Medicine name:
Nafpenzal DC intramamálna suspenzia pre hovädzí dobytok (kravy v období státia na sucho)

Active substance:
Dihydrostreptomycin

Nafcillin

Benzylpenicillin procaine

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Dihydrostreptomycin
100.00 milligram(s) / 1.00 Syringe

Nafcillin
100.00 milligram(s) / 1.00 Syringe

Benzylpenicillin procaine
300000.00 international unit(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (dry cow)**
  - Meat and offal. 14 day
  - Milk. no withdrawal period

36 hours after calving. The product must be applied at least 42 days before calving. If the cow calves before the end of the 42-day period, the remaining days must be added to the 36-hour withdrawal period for milk. At the same time, in such a case, it is necessary to check the presence of residues in the milk.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
29/04/1992

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/330/92-S

Date of authorisation status change:
29/04/1992

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.