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Nafpenzal DC suspensija ievadīšanai tesmenī liellopiem

Not authorised

- Benzylpenicillin procaine
- Dihydrostreptomycin sulfate
- Nafcillin sodium

Product identification

Medicine name:
Nafpenzal DC suspensija ievadīšanai tesmenī liellopiem

Active substance:
Benzylpenicillin procaine

Dihydrostreptomycin sulfate

Nafcillin sodium

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Benzylpenicillin procaine

300.00 milligram(s) / 1.00 Syringe

Dihydrostreptomycin sulfate
100.00 milligram(s) / 1.00 Syringe

Nafcillin sodium
100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (dry cow)**
  - Meat and offal. 14 day

Pienam: ārstēšanas-dzemdību intervāls ≥ 42 dienas: 36 stundas pēc dzemdībām ārstēšanas-dzemdību intervāls < 42 dienas: piens jāpārbauda uz antibiotiku atliekām pirms piegādes.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

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

Authorisation status:
Surrendered

Authorised in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
14/10/1993

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

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/93/0034

Date of authorisation status change:
16/09/2026

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

Documents

Combined File of all Documents

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