Vanaproc 333 mg/g – Suspension zur intramammären Anwendung für Rinder

Authorised

- Benzylpenicillin procaine monohydrate

Product identification

Medicine name:
Vanaproc 333 mg/g – Suspension zur intramammären Anwendung für Rinder

Active substance:
Benzylpenicillin procaine monohydrate

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Benzylpenicillin procaine monohydrate
3.00 gram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (lactating cow)**

  - Meat and offal. 5 day

1 Euterinjektor (3 g Procain-Benzylpenicillin 1 H2O) pro Euterviertel

  - Milk. 120 hour

1 Euterinjektor (3 g Procain-Benzylpenicillin 1 H2O) pro Euterviertel

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51CE09

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

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vana Ges.m.b.H.

Marketing authorisation date:
6/04/1992

Manufacturing sites for batch release:
Vana Ges.m.b.H.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00169

Date of authorisation status change:
6/04/1992

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

Documents

Package Leaflet

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Summary of Product Characteristics

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

Labelling

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