Terramicina 500 mg comprimidos intrauterinos

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:
Terramicina 500 mg comprimidos intrauterinos

Active substance:
Oxytetracycline hydrochloride

Target species:
Cattle (cow for reproduction)

Route of administration:
Intrauterine use

Product details

Active substance and strength:
Oxytetracycline hydrochloride
500.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Intrauterine tablet

Withdrawal period by route of administration:

Intrauterine use:

- **Cattle (cow for reproduction)**
  - Meat and offal. 4 day
  - Milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AA01

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

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Available only in Portuguese
Available only in Portuguese

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis Portugal Lda.

Marketing authorisation date:
4/02/1980

Manufacturing sites for batch release:
Farmasierra Laboratorios S.L.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1468/01/21NFVPT

Date of authorisation status change:
1/11/2021

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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