FINOXALINE

Not authorised

- Flunixin meglumine
- Oxytetracycline hydrochloride

Product identification

Medicine name:
FINOXALINE

Active substance:
Flunixin meglumine

Oxytetracycline hydrochloride

Target species:
Cattle

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Flunixin meglumine
33.20 milligram(s) / 1.00 millilitre(s)

Oxytetracycline hydrochloride
108.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 39 day
  - Milk. 7 day

Intravenous use:

- **Cattle**
  - Meat and offal. 39 day
  - Milk. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA56

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

Authorisation status:
Surrendered

Authorised in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Intervet

Marketing authorisation date:
10/12/1985

Manufacturing sites for batch release:
Trirx Segre

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/3987603 2/1985

Date of authorisation status change:
4/07/2023

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

Documents

Summary of Product Characteristics

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