Ceporex 180 mg/ml Suspension for Injection for cattle. cats, and dogs

Authorised

- Cefalexin sodium

Product identification

Medicine name:
Ceporex 180 mg/ml Suspension for Injection for cattle. cats, and dogs

Active substance:
Cefalexin sodium

Target species:
Cattle
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Cefalexin sodium

19.14 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 15 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01DB01

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
Colourless, multidose 100ml Type I or Type II glass vial, sealed with a bromobutyl rubber closure and an aluminium cap with tear-off lid

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Intervet (Ireland) Limited

Marketing authorisation date:
1/10/1998

Manufacturing sites for batch release:
Aprilia Animal Health S.r.l.
Intervet International GmbH

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/223/001

Date of authorisation status change:
1/10/1998

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

Documents

Summary of Product Characteristics