Naxcel 200 mg/ml - Suspension for injection

Authorised

- Ceftiofur

Product identification

Medicine name:
Naxcel 200 mg/ml - Suspension for injection

Active substance:
Ceftiofur

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Ceftiofur
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 9 day 9 days
  - Milk. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Cyprus , Czechia , Estonia , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Poland , Slovakia ,

United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:1 vial, Content:100 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
19/05/2005

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
8/10/2009

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

Documents

Combined File of all Documents

English (PDF)
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