Convenia 80 mg/ml - Powder and solvent for solution for injection

Authorised

- Cefovecin

Product identification

Medicine name:
Convenia 80 mg/ml - Powder and solvent for solution for injection

Active substance:
Cefovecin

Target species:
Dog
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Cefovecin
80.00 milligram(s) / 1.00 Vial

Pharmaceutical form:
Powder and solvent for solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01DD91

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

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:
Belgium , Bulgaria , Cyprus , Czechia , Denmark , Estonia , Finland , Greece , Hungary , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Powder: vial (glass), solvent: vial (glass), Package_size:1 vial + 1 vial, Content:Powder: 340 mg, solvent: 4.45 ml
Packaging:Powder: vial (glass), solvent: vial (glass), Package_size:1 vial + 1 vial, Content:Powder: 852 mg, solvent: 10.8 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 SA

Marketing authorisation date:
19/06/2006

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.

Responsible authority:
European Commission

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

Date of authorisation status change:
19/06/2006

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

Documents

Combined File of all Documents

English (PDF)
Published on: 20/06/2024
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