

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

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:

Austria , 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

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)

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