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Virbagen Omega (--) - Powder and solvent for suspension for injection

Authorised

- Interferon omega, recombinant, feline

Product identification

Medicine name:
Virbagen Omega (--) - Powder and solvent for suspension for injection

Active substance:
Interferon omega, recombinant, feline

Target species:
Dog
Cat

Route of administration:
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Interferon omega, recombinant, feline
Presentation_strength:10 MU Reference:Hse Index:0

Pharmaceutical form:
Powder and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QL03AB

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 , Croatia , Cyprus , Czechia , Estonia , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , United Kingdom (Northern Ireland)

Package description:
Packaging:Powder: vial (glass); solvent: vial (glass), Package_size:5 vials (Powder) + 5 vials (Solvent), Content:10 MU (Powder); 1 ml (Solvent)
Packaging:Powder: vial (glass); solvent: vial (glass), Package_size:2 vials (Powder) + 2 vials (Solvent), Content:10 MU (Powder) ; 1 ml (Solvent)
Packaging:Powder: vial (glass); solvent: vial (glass), Package_size:1 vial (Powder) + 1 vial (Solvent), Content:10 MU (Powder) ; 1 ml (Solvent)

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:

Virbac

Marketing authorisation date:
6/11/2001

Manufacturing sites for batch release:
Virbac

Responsible authority:
European Commission

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

Date of authorisation status change:
6/11/2001

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: 14/03/2026
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