

Imrestor 5.5 mg/ml - Solution for injection in pre-filled syringe

Authorised

- Pegbovigrastim

Product identification

Medicine name:

Imrestor 5.5 mg/ml - Solution for injection in pre-filled syringe

Active substance:

Pegbovigrastim

Target species:

Cattle (heifer)

Cattle (dairy cow)

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Pegbovigrastim

5.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QL03AA90

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)

Package description:

Packaging:Pre -filled syringe (PP), Package_size:10 pre-filled syringes, Content:2.7 ml

Packaging:Pre-filled syringe (PP), Package_size:50 pre-filled syringes, Content:2.7 ml

Packaging:Pre -filled syringe (PP), Package_size:100 pre-filled syringes, Content:2.7 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Elanco GmbH

Marketing authorisation date:

9/12/2015

Manufacturing sites for batch release:

Elanco France S.A.S

TriRx Speke Limited

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

9/12/2015

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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