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Improvac (--) - Solution for injection

Authorised

- Gonadotropin releasing factor analogue diphtheria toxoid conjugate

Product identification

Medicine name:
Improvac (--) - Solution for injection

Active substance:
Gonadotropin releasing factor analogue diphtheria toxoid conjugate

Target species:
Pig

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Gonadotropin releasing factor analogue diphtheria toxoid conjugate
Presentation_strength:300 µg Reference:Hse Index:0

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Pig**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG03XA91

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 , Czechia , Denmark , Finland , Germany , Greece , Hungary , Iceland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Polyethylene bottle with rubber closure and aluminium cap,
Package_size:10 bottles, Content:100 ml
Packaging:Polyethylene bottle with rubber closure and aluminium cap,
Package_size:4 bottles, Content:250 ml
Packaging:Polyethylene bottle with rubber closure and aluminium cap,
Package_size:1 bottle, Content:100 ml
Packaging:Polyethylene bottle with rubber closure and aluminium cap,
Package_size:1 bottle, Content:250 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
11/05/2009

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:
11/05/2009

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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