OESTRACTON 52.4 micrograms/ml solution for injection for cattle, horses, pigs

Authorised

- Gonadorelin (6-D-phenylalanine) acetate

Product identification

Medicine name:
OESTRACTON 52.4 micrograms/ml solution for injection for cattle, horses, pigs

Active substance:
Gonadorelin (6-D-phenylalanine) acetate

Target species:
Pig
Cattle
Horse

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Gonadorelin (6-D-phenylalanine) acetate

52.40 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Pig**
  - Meat and offal. 0 day

Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01CA01

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

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (glass) with 50 millilitre(s)
(ID2) 60 millilitre(s): unspecified outer container with 6 Vial (glass) each with 10 millilitre(s)
(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (glass) with 10 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol

Marketing authorisation date:
5/11/2013

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V444841

Date of authorisation status change:
5/11/2013

Reference member state:
Germany

Procedure number:
DE/V/0154/001

Concerned member states:

Belgium Czechia France Hungary Ireland Netherlands
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (RTF)
Published on: 16/02/2026
Download

Package Leaflet

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Labelling

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