

Oestracton 52.4 microgram/ml solution for injection for cattle, horses, pigs

Authorised

- Gonadorelin (6-D-phenylalanine) acetate

Product identification

Medicine name:

Oestracton 52.4 microgram/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:

Ireland

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 S.A.

Marketing authorisation date:

8/11/2013

Manufacturing sites for batch release:

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10521/003/001

Date of authorisation status change:

8/11/2013

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

2401818-paren-20140220.pdf