

Folligon 200 IU/ml lyofilizát na injekčný roztok

Not authorised

- Serum gonadotrophin
- Serum gonadotrophin

Product identification

Medicine name:

Folligon 200 IU/ml lyofilizát na injekčný roztok

Active substance:

Serum gonadotrophin

Serum gonadotrophin

Target species:

Cattle (cow)

Cattle (heifer)

Sheep (ewe)

Goat (adult female)

Pig (sow)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Serum gonadotrophin
5000.00 international unit(s) / 1.00 Vial

Serum gonadotrophin
1000.00 international unit(s) / 1.00 Vial

Pharmaceutical form:

Lyophilisate and solvent for solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle (cow)

- Meat and offal. 0 day Zero days.

•

Cattle (heifer)

- Meat and offal. 0 day Zero days.

•

Sheep (ewe)

- Meat and offal. 0 day Zero days.

•

Goat (adult female)

- Meat and offal. 0 day Zero days.

•

Pig (sow)

- Meat and offal. 0 day Zero days.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG03GA03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Slovakia

Package description:

Available only in [Slovak](#)

Available only in [Slovak](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

13/10/2010

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/044/10-S

Date of authorisation status change:

20/01/2025

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

Documents

Combined File of all Documents

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