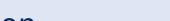
Folligon PMSG 200 IU/ml lyophilisate and solvent for solution for injection



Gonadotropin, equine, serum

Product identification

Medicine name: Folligon PMSG 200 IU/ml lyophilisate and solvent for solution for injection

Active substance:

Gonadotropin, equine, serum

Target species:

Cattle Sheep Pig

Route of administration:

Intramuscular use Subcutaneous use

Product details

Active substance and strength:

Gonadotropin, equine, serum 5000.00 international unit(s) / 1.00 Vial Authorised

Pharmaceutical form:

Lyophilisate and solvent for solution for injection

Withdrawal period by route of administration:

Intramuscular use:

Cattle

- Meat and offal. 0 day
- Milk. 0 day

Sheep

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- Meat and offal. 0 day

- Milk. 0 day

Pig

- Meat and offal. 0 day

Subcutaneous use:

Pig

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- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG03GA03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Lyophilisate: Clear, type I (Ph. Eur.) glass vials with halogenobutyl rubber stopper, closed with a colour codedaluminium cap.Solvent: Clear, type I (Ph. Eur) or type II (Ph. Eur.) glass vials with halogenobutyl rubber stopper, closed with a colourcoded aluminium cap.Pack sizes:Cardboard box with 1 vial of 1,000 IU lyophilisate and 1 x 5 ml vial of solvent.

Lyophilisate: Clear, type I (Ph. Eur.) glass vials with halogenobutyl rubber stopper, closed with a colour codedaluminium cap.Solvent: Clear, type I (Ph. Eur) or type II (Ph. Eur.) glass vials with halogenobutyl rubber stopper, closed with a colourcoded aluminium cap.Pack sizes:Cardboard box with 1 vial of 5,000 IU lyophilisate and 1 x 25 ml vial of solvent.

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:

Intervet (Ireland) Limited

Marketing authorisation date:

19/09/1998

Manufacturing sites for batch release:

Intervet International GmbH Intervet International B.V.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10996/055/001

Date of authorisation status change:

19/09/1998

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

Documents

Summary of Product Characteristics

Source URL: https://medicines.health.europa.eu/veterinary/60000089489