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

Authorised

- 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

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

Subcutaneous use:

- **Pig**
  - 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
