Folltropin 700 IU Powder and Solvent for solution for injection

Authorised

- Follicle stimulating hormone, porcine
- Water for injection

Product identification

Medicine name:
Folltropin 700 IU Powder and Solvent for solution for injection
FOLLTROPIN 700 UI POUDRE ET SOLVANT POUR SOLUTION INJECTABLE

Active substance:
Follicle stimulating hormone, porcine

Water for injection

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Follicle stimulating hormone, porcine
700.00 international unit(s) / 1.00 Vial

Water for injection
1.00 other / 1.00 Vial

Pharmaceutical form:
Powder and solvent for solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG03GA90

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Cardboard box containing one vial of powder and one vial of solvent.Freeze-dried powderClear glass 20 ml vial (Type I), with halobutyl rubber stopper (Type I) and red flip-off cap.SolventClear glass 20 ml vial (Type I), with halobutyl rubber stopper (Type I) and yellow flip-off cap.

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:
Vetoquinol S.A.

Marketing authorisation date:
26/06/2013

Manufacturing sites for batch release:
Vetoquinol

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/3692745 0/2013

Date of authorisation status change:
21/07/2017

Reference member state:
Ireland

Procedure number:
IE/V/0126/001

Concerned member states:
Austria Czechia Denmark France Germany Italy Netherlands Poland Spain 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 (PDF)
Published on: 25/09/2024

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Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents
