PLUSET 500 IU/500 IU powder and solvent for solution for injection for cattle

Authorised

- FOLLICLE-STIMULATING HORMONE
- Luteinising hormone

Product identification

Medicine name:
PLUSET 500 IU/500 IU powder and solvent for solution for injection for cattle
Pluset Vet. 500+500 IE pulver og solvens til injektionsvæske, opløsning

Active substance:
FOLLICLE-STIMULATING HORMONE
Luteinising hormone

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
FOLLICLE-STIMULATING HORMONE
500.00 international unit(s) / 10.00 millilitre(s)

Luteinising hormone
500.00 international unit(s) / 10.00 millilitre(s)

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:
Denmark

Available in:
Denmark

Package description:
Cardboard box with 2 glass vials of 10 ml of lyophilised product and 1 glass vial of 21 ml of diluent. Lyophilised product: Vial of colourless neutral glass (type 1) Capacity: 10 ml. Provided with bromobutyl and silicate stopper and aluminium cap flip off seal. Container for the diluent: Vial of colourless neutral glass (type 1) Capacity: 21 ml. With rubber penitype stopper of grey colour and aluminium cap flip off seal.

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:
Laboratorios Calier S.A.

Marketing authorisation date:
8/09/2008

Manufacturing sites for batch release:
Laboratorios Calier S.A.

Responsible authority:
Danish Medicines Agency

Authorisation number:
42390

Date of authorisation status change:
8/09/2008

Reference member state:
Italy

Procedure number:
IT/V/0117/001

Concerned member states:
Belgium Czechia Denmark Finland France Germany Greece Hungary Ireland Luxembourg Netherlands Poland Portugal 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

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