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

Authorised

- Luteinising hormone
- FOLLICLE-STIMULATING HORMONE

## Product identification

**Medicine name:**
PLUSET 500 IU/500 IU powder and solvent for solution for injection for cattle
Pluset poeder en oplosmiddel voor oplossing voor injectie

**Active substance:**
Luteinising hormone
FOLLICLE-STIMULATING HORMONE

**Target species:**
Cattle

**Route of administration:**
Intramuscular use

## Product details

**Active substance and strength:**
Luteinising hormone

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

FOLLICLE-STIMULATING 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:**
Netherlands

**Available in:**
Netherlands

**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:**
16/06/2004

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

**Responsible authority:**
Medicines Evaluation Board

**Authorisation number:**
REG NL 10171

**Date of authorisation status change:**
7/07/2022

**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

Combined File of all Documents

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