

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

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.

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

English (PDF)

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Summary of Product Characteristics