VETECORH 1000 IU/ML LYOPHILISATE AND SOLVENT FOR SOLUTION FOR INJECTION FOR CATTLE, HORSES, SHEEP, GOATS, PIGS, CATS AND DOGS

Authorised

- HUMAN CHORIONIC GONADOTROPIN

Product identification

Medicine name:
VETECORH 1000 IU/ML LYOPHILISATE AND SOLVENT FOR SOLUTION FOR INJECTION FOR CATTLE, HORSES, SHEEP, GOATS, PIGS, CATS AND DOGS

Active substance:
HUMAN CHORIONIC GONADOTROPIN

Target species:
Cattle
Pig
Cat
Horse
Sheep
Goat
Dog

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
HUMAN CHORIONIC GONADOTROPIN
5000.00 international unit(s) / 1.00 Bottle

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
- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**

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

Intravenous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG03GA01

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

---

Authorisation status:
Valid

Authorised in:
Poland

Package description:
Box of 1 vial of lyophilisate + Box of 1 vial of 5 ml of solvent
Box of 2 vials of lyophilisate + Box of 2 vials of 5 ml of solvent
Box of 5 vials of lyophilisate + Box of 5 vials of 5 ml of solvent
Box of 1 vial of lyophilisate and 1 vial 5 ml of solvent
Box of 2 vials of lyophilisate and 2 vials 5 ml of solvent
Box of 5 vials of lyophilisate and 5 vials of 5 ml of solvent

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application - change in active substance(s) (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Laboratorios Calier S.A.

Marketing authorisation date:
17/06/2025

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3420

Date of authorisation status change:
17/06/2025

Reference member state:

France

Procedure number:
FR/V/0467/001

Concerned member states:
Austria Belgium Bulgaria Croatia Germany Hungary Ireland Italy Latvia Lithuania Netherlands Poland Portugal Romania 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

eu-puar-frv0467001-mr-rpe883-en.pdf