Tylolab tartrate 200,000 IU/ml solution for injection

Authorised

- Tylosin tartrate

Product identification

Medicine name:
Tylolab tartrate 200,000 IU/ml solution for injection
Tylolab tartrate 200.000 IU/ml ενέσιμο διάλυμα

Active substance:
Tylosin tartrate

Target species:
Cattle
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Tylosin tartrate
200000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 33 day
  - Milk. 120 hour
- **Pig**
  - Meat and offal. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

Amber type II glass vials containing 250 mL veterinary medicinal product, closed with a chlorobutyl stopper and aluminium caps.
Amber type II glass vials containing 250 mL veterinary medicinal product, closed with a chlorobutyl stopper and aluminium caps.
Amber type II glass vials containing 250 mL veterinary medicinal product, closed with a chlorobutyl stopper and aluminium caps.
Amber type II glass vials containing 100 mL veterinary medicinal product, closed with a chlorobutyl stopper and aluminium caps.

Amber type II glass vials containing 100 mL veterinary medicinal product, closed with a chlorobutyl stopper and aluminium caps.

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:
Labiana Life Sciences S.A.

Marketing authorisation date:
12/06/2023

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
62551/13.06.2023/27-07-2023/K-0252201

Date of authorisation status change:
26/07/2023

Reference member state:
Hungary

Procedure number:
HU/V/0147/001

Concerned member states:
Cyprus France Greece Ireland Romania United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet