

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 NE/ml oldatos injekció

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:

Hungary

Available in:

Hungary

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:

6/06/2023

Manufacturing sites for batch release:

LABIANA LIFE SCIENCES, S.A.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

6/06/2023

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