

Labiprofen 150 mg/ml solution for injection

Authorised

- Ketoprofen

Product identification

Medicine name:

Labiprofen 150 mg/ml solution for injection

Labiprofen 150 mg/ml oldatos injekció szarvasmarhák, sertések és lovak részére
A.U.V.

Active substance:

Ketoprofen

Target species:

Cattle

Pig

Horse

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Ketoprofen

150.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:****• Cattle**

- Meat and offal. 2 day

• Pig

- Meat and offal. 3 day

• Cattle

- Milk. 0 hour

• Horse

- Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

Intravenous use:**• Cattle**

- Meat and offal. 2 day

• Horse

- Meat and offal. 1 day

• Cattle

- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AE03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Package description:

box containing 10 vials of 250 ml

box containing 10 vials of 100 ml
box containing 12 vials of 50 ml
box containing 1 vial of 250 ml
box containing 1 vial of 100 ml
box containing 1 vial of 50 ml

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:

20/01/2021

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

4229/X/21 NÉBIH ÁTI

Date of authorisation status change:

20/01/2021

Reference member state:

Spain

Procedure number:

ES/V/0388/001

Concerned member states:

Austria Belgium Bulgaria Croatia Denmark Estonia Finland France Germany
Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Norway Portugal
Romania Slovenia Sweden United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

Labelling

Summary of Product Characteristics

Source URL: <https://medicines.health.europa.eu/veterinary/600000017650>