KALIPROFEN LA 50 mg/ml raztopina za injiciranje za govedo

Authorised

- Carprofen

Product identification

Medicine name:
KALIPROFEN LA 50 mg/ml raztopina za injiciranje za govedo

Active substance:
Carprofen

Target species:
Cattle

Route of administration:
Subcutaneous use
Intravenous use

Product details

Active substance and strength:
Carprofen
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 21 day — Meso in organi: 21 dni
  - Milk. 0 hour — Mleko: nič ur

Intravenous use:

- **Cattle**
  - Milk. 0 hour — Mleko: nič ur
  - Meat and offal. 21 day — Meso in organi: 21 dni

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AE91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Slovenia

Package description:

Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
GENERA SI podjetje za zastopanje in trgovino d.o.o.

Marketing authorisation date:
28/11/2021

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
NP/V/0418/001

Date of authorisation status change:
14/12/2012

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.