REUFLOGIN, 50mg/ml, Injekční roztok

Authorised

- Diclofenac sodium

Product identification

Medicine name:
REUFLOGIN, 50mg/ml, Injekční roztok

Active substance:
Diclofenac sodium

Target species:
Horse

Route of administration:
Intravenous use
Intramuscular use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse**

- Meat and offal. no withdrawal period

Nepoužívat u koní, jejichž maso je určeno pro lidskou spotřebu.,

- Milk. no withdrawal period

Nepoužívat u koní, jejichž mléko je určeno pro lidskou spotřebu.,

Intramuscular use:

- **Horse**

- Meat and offal. no withdrawal period

Nepoužívat u koní, jejichž maso je určeno pro lidskou spotřebu.,

- Milk. no withdrawal period

Nepoužívat u koní, jejichž mléko je určeno pro lidskou spotřebu.,

---

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AB05

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

---

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
19/09/1997

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/844/97-C

Date of authorisation status change:
19/09/1997

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

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

Labelling

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