

Niglumine, 50mg/ml, Injekční roztok

Authorised

- Flunixin

Product identification

Medicine name:

Niglumine, 50mg/ml, Injekční roztok

Active substance:

Flunixin

Target species:

Horse

Cattle

Route of administration:

Intravenous use

Product details

Active substance and strength:

Flunixin

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intravenous use:**

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Horse

- Meat and offal. no withdrawal period

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

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Cattle

- Meat and offal. 7 day

- Milk. 48 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

Available only in [Czech](#)

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:

Laboratorios Calier S.A.

Marketing authorisation date:

20/02/2002

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/023/02-C

Date of authorisation status change:

29/11/2010

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.

Labelling

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.