

FLUNEX 50 MG / ML SOLUTION FOR INJECTION FOR CATTLE, HORSES AND PIGS

Authorised

- Flunixin meglumine

Product identification

Medicine name:

FLUNEX 50 MG / ML SOLUTION FOR INJECTION FOR CATTLE, HORSES AND PIGS

Active substance:

Flunixin meglumine

Target species:

Cattle

Pig

Horse

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Flunixin meglumine

82.90 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

- Meat and offal. 31 day

- Milk. 36 hour

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Pig

- Meat and offal. 24 day

Intravenous use:

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Cattle

- Meat and offal. 4 day

- Milk. 24 hour

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Horse

- Milk. no withdrawal period

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

- Meat and offal. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

1x50 ml vial

1x100 ml vial

1x250 ml vial

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Industrial Veterinaria S.A.

Marketing authorisation date:

7/11/2024

Manufacturing sites for batch release:

aniMedica GmbH

Industrial Veterinaria S.A.

aniMedica Herstellungs GmbH

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/057/24-C

Date of authorisation status change:

7/11/2024

Reference member state:

France

Procedure number:

FR/V/0474/001

Concerned member states:

Austria Cyprus Czechia Estonia Germany Greece Hungary Ireland Italy
Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Spain
United Kingdom (Northern Ireland)

Generic of:

600000038769

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.

Combined File of all Documents

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