**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000061270

# Flunixin 50 mg/ml Solution for Injection for Cattle, Horses and Pigs

Authorised

Flunixin meglumine

# Product identification

#### **Medicine name:**

Flunixin 50 mg/ml Solution for Injection for Cattle, Horses and Pigs Flunixin N-vet 50 mg/ml Injektionsvätska, lösning

#### **Active substance:**

Flunixin meglumine

# **Target species:**

Cattle

Horse

Pig

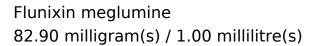
#### **Route of administration:**

Intravenous use

Intramuscular use

# **Product details**

# **Active substance and strength:**



#### **Pharmaceutical form:**

Solution for injection

# Withdrawal period by route of administration: Intravenous use:

Cattle

- Milk. 36 hour
- Meat and offal. 7 day

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#### Horse

- Milk. no withdrawal period

Do not use in mares producing milk for human consumption.

- Meat and offal. 7 day

#### Intramuscular use:

Pig

- Meat and offal. 22 day

# Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG90

# Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Sweden

#### **Available in:**

#### Sweden

#### Package description:

Available only in Swedish

Available only in Swedish

Available only in Swedish

Available only in **Swedish** 

Available only in Swedish

Available only in **Swedish** 

Available only in Swedish

Available only in Swedish

Available only in Swedish

Available only in **Swedish** 

### 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:

Norbrook Laboratories (Ireland) Limited

# Marketing authorisation date:

27/10/2009

# Manufacturing sites for batch release:

Norbrook Laboratories (Ireland) Limited

Norbrook Laboratories Limited

# Responsible authority:

Swedish Medical Products Agency

#### **Authorisation number:**

27763

# Date of authorisation status change:

R	efe	ran	6	m	am	har	cta	ta:
П	CIC	CII	CC	1117		NCI	3 LC	ILC.

Germany

#### **Procedure number:**

DE/V/0325/001

#### **Concerned member states:**

Iceland Netherlands Portugal Sweden United Kingdom (Northern Ireland)

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

## **Documents**

Package Leaflet

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

Summary of Product Characteristics

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