

Καλαφεν 50 mg/ml, ενέσιμο διάλυμα για βοοειδή, άλογα και χοίρους

Authorised

- Flunixin meglumine

Product identification

Medicine name:

Kalafev 50 mg/ml, ενέσιμο διάλυμα για βοοειδή, άλογα και χοίρους

Active substance:

Flunixin meglumine

Target species:

Cattle

Horse

Pig

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Flunixin meglumine

82.95 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

- Meat and offal. 5 day

Δεν επιτρέπεται η χορήγηση σε ζώα τα οποία παράγουν γάλα για ανθρώπινη κατανάλωση

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

Available only in [Greek](#)

Available only in [Greek](#)

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic (abridged application) - art 13(1)

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

7/04/2013

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

69428/18/26-11-2019/K-0186601

Date of authorisation status change:

7/09/2021

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

Documents

Summary of Product Characteristics

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Package Leaflet

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