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GENIXINE

Authorised

- Flunixin meglumine

Product identification

Medicine name:

GENIXINE

Active substance:

Flunixin meglumine

Target species:

Cattle

Pig

Horse

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Flunixin meglumine

83.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

•

Cattle

- Meat and offal. 31 day
- Milk. 36 hour

•

Pig

- Meat and offal. 20 day

Intravenous use:

•

Cattle

- Meat and offal. 10 day
- Milk. 24 hour

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Horse

- Meat and offal. 10 day
- Milk. no withdrawal period

Lait : ne pas utiliser chez les juments en lactation productrices de lait destiné à la consommation humaine

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in [French](#)

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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:

Ceva Sante Animale

Marketing authorisation date:

12/03/2010

Manufacturing sites for batch release:

Ceva Sante Animale

Vetem S.p.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/3839789 3/2010

Date of authorisation status change:

12/03/2015

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 and Labelling

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