

DINOLYTIC 12,5 MG/ML SOLUTION INJECTABLE POUR BOVINS

Not
authorised

- Dinoprost

Product identification

Medicine name:

DINOLYTIC 12,5 MG/ML SOLUTION INJECTABLE POUR BOVINS

Active substance:

Dinoprost

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Dinoprost

12.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- Cattle

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Zoetis France

Marketing authorisation date:

4/05/2018

Manufacturing sites for batch release:

Zoetis Manufacturing & Research Spain S.L.

Responsible authority:

Authorisation number:

FR/V/9364757 5/2018

Date of authorisation status change:

23/06/2022

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.

Source URL: <https://medicines.health.europa.eu/veterinary/600000040650>