KETOXYME

Authorised

- Ketoprofen

Product identification

Medicine name:
KETOXYME
Ketoxyme 100 mg/ml oplossing voor gebruik in drinkwater

Active substance:
Ketoprofen

Target species:
Pig

Route of administration:
In drinking water use

Product details

Active substance and strength:
Ketoprofen
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **Pig**

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
bottle of 1 L

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:
Andersen S.A.

Marketing authorisation date:
This information is not available for this product.

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 114768

Date of authorisation status change:
18/01/2022

Reference member state:
Spain

Procedure number:
ES/V/0213/001

Concerned member states:
Belgium Netherlands United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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

Summary of Product Characteristics

Package Leaflet

eu-PUAR-ketoxyme-en.pdf
