

Pyrocam 15 mg/ml oral suspension for pigs

Authorised

- Meloxicam

Product identification

Medicine name:

Pyrocam 15 mg/ml oral suspension for pigs

Pyrocam 15 mg/ml suspensão oral para suínos

Active substance:

Meloxicam

Target species:

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Meloxicam

15.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:**Oral use:**

-

Pig

- Meat and offal. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AC06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Package description:

Cardboard box with HDPE bottle of 1000 ml closed with a two-part tamper evident closure consisting of an outer white cap (PP), an internal natural colour screw closure (HDPE) and a mounted plug (LDPE) + plastic measuring syringe

Cardboard box with HDPE bottle of 250 ml closed with a two-part tamper evident closure consisting of an outer white cap (PP), an internal natural colour screw closure (HDPE) and a mounted plug (LDPE) + plastic measuring syringe

Cardboard box with HDPE bottle of 125 ml closed with a two-part tamper evident closure consisting of an outer white cap (PP), an internal natural colour screw closure (HDPE) and a mounted plug (LDPE) + plastic measuring syringe

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application – bioavailability studies cannot be used to demonstrate bioequivalence (Article 19(1)(b) of Regulation (EU) 2019/6)

Marketing authorisation holder:

HuVepharma

Marketing authorisation date:

20/10/2023

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

600/01/23DFVPT

Date of authorisation status change:

20/10/2023

Reference member state:

Belgium

Procedure number:

BE/V/0048/001

Concerned member states:

Austria Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France
Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain
Sweden 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

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