# Melovem 20 mg/ml - Solution for injection (Horse, Cattle, Pig)

Authorised

Meloxicam

# Product identification

#### **Medicine name:**

Melovem 20 mg/ml - Solution for injection (Horse, Cattle, Pig)

#### **Active substance:**

Meloxicam

#### **Target species:**

Cattle

Horse

Pig

#### **Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

# **Product details**

### **Active substance and strength:**

Meloxicam

20.00 milligram(s) / 1.00 millilitre(s)

#### **Pharmaceutical form:**

# Withdrawal period by route of administration: Intramuscular use:

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#### Cattle

- Meat and offal. 15 day 15 days
- Milk. 5 day 5 days

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#### Horse

- Meat and offal. 5 day

5 days. Not authorised to use in horses producing milk for human consumption.

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# Pig

- Meat and offal. 5 day 5 days

# Intravenous use:

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#### Cattle

- Meat and offal. 15 day 15 days
- Milk. 5 day 5 days

•

#### Horse

- Meat and offal. 5 day

5 days. Not authorised to use in horses producing milk for human consumption.

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#### Pig

- Meat and offal. 5 day 5 days

#### Subcutaneous use:

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#### Cattle

- Meat and offal. 15 day 15 days
- Milk. 5 day 5 days

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#### Horse

- Meat and offal. 5 day

5 days. Not authorised to use in horses producing milk for human consumption.

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# Pig

- Meat and offal. 5 day 5 days

# Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AC06

# Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### Authorised in:

Austria, Belgium, Bulgaria, Croatia, Cyprus, Czechia, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, United Kingdom (Northern Ireland)

# **Package description:**

Packaging:Vial (glass), Package\_size:1 vial, Content:250 ml Packaging:Vial (glass), Package\_size:1 vial, Content:100 ml Packaging:Vial (glass), Package size:1 vial, Content:50 ml

# Additional information

Entitlement type:
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Marketing Authorisation

#### Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

#### Marketing authorisation holder:

Dopharma Research B.V.

#### Marketing authorisation date:

7/07/2009

# Manufacturing sites for batch release:

Dopharma B.V.

#### **Responsible authority:**

**European Commission** 

#### **Authorisation number:**

This information is not available for this product.

# **Date of authorisation status change:**

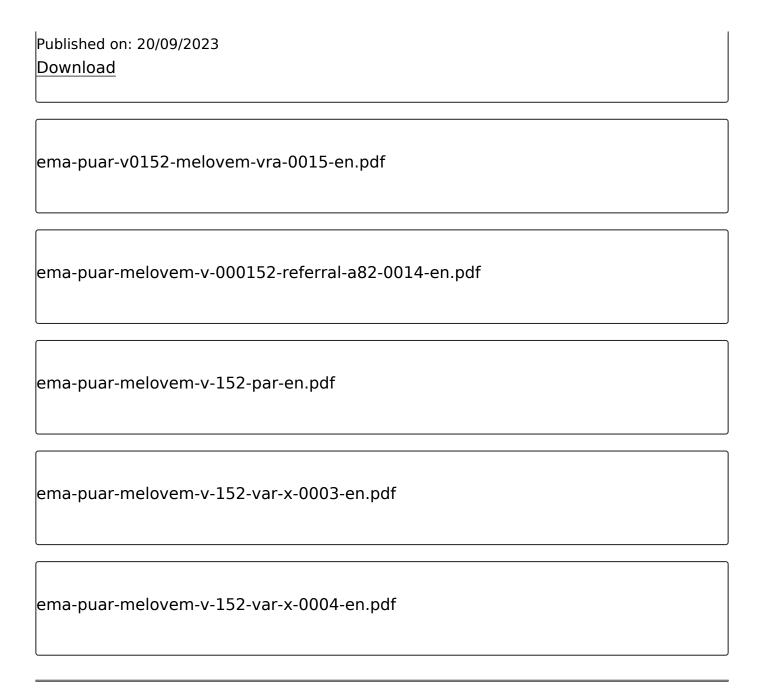
25/09/2013

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

# **Documents**

Combined File of all Documents

English (PDF)



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