Source URL: https://medicines.health.europa.eu/veterinary/en/600000984270

Metaxx 5 mg/ml solution for injection for cattle, pigs, dogs and cats

Authorised

Meloxicam

Product identification

Medicine name:

Metaxx 5 mg/ml solution for injection for cattle, pigs, dogs and cats Mexxam Vet 5 mg/ml šķīdums injekcijām liellopiem, cūkām, suņiem un kaķiem

Active substance:

Meloxicam

Target species:

Dog

Cat

Cattle

Pig

Route of administration:

Subcutaneous use

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Meloxicam

5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Concentrate for solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

Dog

- Not applicable. no withdrawal period

Cat

- Not applicable. no withdrawal period

Cattle

- Meat and offal. no withdrawal period 15 days

Intramuscular use:

Pig

- Meat and offal. no withdrawal period 5 days

Intravenous use:

Cattle

- Meat and offal. no withdrawal period

15 days Not authorised for use in animals producing milk for human consumption.

Legal status of supply: Veterinary medicinal product subject to veterinary prescription	
Authorisation status: Valid	
Authorised in: Latvia	
Package description:	
Available only in <u>Latvian</u>	
Additional information	
Entitlement type:	
Marketing Authorisation	
Legal basis of product authorisation: Generic (abridged application) - art 13(1)	
Marketing authorisation holder: Alfasan Nederland B.V.	
Marketing authorisation date:	

Responsible authority:

Alfasan Nederland B.V.

Food And Veterinary Service

Authorisation number:

V/DCP/22/0046

Date of authorisation status change:

30/11/2022

Reference member state:

Netherlands

Procedure number:

NL/V/0375/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg 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

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