Hyaluronan 10 mg/ml Roztwór do wstrzykiwań

Authorised

This information is not available for this product.

Product identification

Medicine name:
Hyaluronan 10 mg/ml Roztwór do wstrzykiwań

Active substance:
This information is not available for this product.

Target species:
Cat
Dog
Horse

Route of administration:
Intraocular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
This information is not available for this product.

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intraocular use:

- **Cat**

  - All relevant tissues. no withdrawal period

The withdrawal period does not apply.

- **Dog**

  - All relevant tissues. no withdrawal period

The withdrawal period does not apply.

- **Horse**

Intravenous use:

- **Cat**

  - All relevant tissues. no withdrawal period

The withdrawal period does not apply.

- **Dog**

  - All relevant tissues. no withdrawal period

The withdrawal period does not apply.

- **Horse**

Subcutaneous use:

- **Cat**

  - All relevant tissues. no withdrawal period

The withdrawal period does not apply.

- **Dog**

  - All relevant tissues. no withdrawal period

The withdrawal period does not apply.

- **Horse**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM09AX01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Grabikowski-Grabikowska Przedsiebiorstwo Produkcyjno-Handlowo-Uslugowe Inex Sp. j.

Marketing authorisation date:
30/01/2017

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2618

Date of authorisation status change:
30/01/2017

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

Documents

Package Leaflet

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

Labelling

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

Summary of Product Characteristics

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