BONHAREN Intravenous, 10mg/ml, Injekční roztok

Authorised

- SODIUM HYALURONATE

Product identification

Medicine name:
BONHAREN Intravenous, 10mg/ml, Injekční roztok

Active substance:
SODIUM HYALURONATE

Target species:
Horse
Cat
Dog

Route of administration:
Intravenous use
Subcutaneous use
Ocular use

Product details

Active substance and strength:
SODIUM HYALURONATE
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour

Subcutaneous use:

- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour

Ocular use:

- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM09AX01

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

Authorisation status:
Valid

Authorised in:
Czechia

Available in:
Czechia

Package description:
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Contipro a.s.

Marketing authorisation date:
8/08/2000

Manufacturing sites for batch release:
Contipro a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/045/00-C

Date of authorisation status change:
2/10/2008

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

Documents

Summary of Product Characteristics

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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.