POLYEQUAN, Injekční suspenze

Authorised

- Immunoglobulins against Escherichia coli, Equine
- Immunoglobulins against Salmonella enterica subsp. enterica, serovar Abortusequi, Equine
- Immunoglobulins against Streptococcus equi, Equine
- Immunoglobulins against Actinobacillus equinum, Equine

Product identification

Medicine name:
POLYEQUAN, Injekční suspenze

Active substance:
Immunoglobulins against Escherichia coli, Equine

Immunoglobulins against Salmonella enterica subsp. enterica, serovar Abortusequi, Equine

Immunoglobulins against Streptococcus equi, Equine

Immunoglobulins against Actinobacillus equinum, Equine

Target species:
Horse (foal)

Route of administration:
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Immunoglobulins against Escherichia coli, Equine
20.00 titre / 1.00 Dose

Immunoglobulins against Salmonella enterica subsp. enterica, serovar Abortusequi, Equine
40.00 titre / 1.00 Dose

Immunoglobulins against Streptococcus equi, Equine
1.00 relative potency / 1.00 Dose

Immunoglobulins against Actinobacillus equinum, Equine
32.00 titre / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse (foal)**
  - Meat and offal. 0 day
  - Milk. 0 hour

Subcutaneous use:

- **Horse (foal)**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AM03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Czechia

Package description:
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:
Bioveta a.s.

Marketing authorisation date:
19/11/1991

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
97/325/91-C

Date of authorisation status change:
18/01/2007

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

Documents

Summary of Product Characteristics

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

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.