Gafervit injekčný roztok

Authorised

- Porcine normal immunoglobulin
- Iron dextran
- Thiamine hydrochloride
- Riboflavin
- Pyridoxine hydrochloride
- Nicotinamide
- Calcium pantothenate
- Copper chloride
- Cobalt(II) chloride

Product identification

Medicine name:
Gafervit injekčný roztok

Active substance:
Porcine normal immunoglobulin
Iron dextran
Thiamine hydrochloride
Riboflavin
Pyridoxine hydrochloride
Nicotinamide
Calcium pantothenate
Copper chloride
Cobalt(II) chloride

Target species:
Pig (piglet)
Pig (suckling piglet)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine normal immunoglobulin
5000.00 milligram(s) / 100.00 millilitre(s)

Iron dextran
700.00 milligram(s) / 100.00 millilitre(s)

Thiamine hydrochloride
3.00 milligram(s) / 100.00 millilitre(s)

Riboflavin
1.14 milligram(s) / 100.00 millilitre(s)

Pyridoxine hydrochloride
0.28 milligram(s) / 100.00 millilitre(s)

Nicotinamide
42.84 milligram(s) / 100.00 millilitre(s)

Calcium pantothenate
1.60 milligram(s) / 100.00 millilitre(s)

Copper chloride
2.71 milligram(s) / 100.00 millilitre(s)

Cobalt(II) chloride
0.27 milligram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig (piglet)**
  - All relevant tissues. 0 day zero days

- **Pig (suckling piglet)**
  - All relevant tissues. 0 day zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB03AE10

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

---

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak

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:
25/07/1994

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
97/396/91-S

Date of authorisation status change:
25/07/1994

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.