Suibiofer Se (586 mg + 0,998 mg + 0,00403 mg + 0,03 mg + 0,01 mg + 0,0025 mg + 0,08 mg + 0,4 mg)/ml, surowica świńska do 1 ml Roztwór do wstrzykiwań

Authorised

- Thiamine hydrochloride
- Pyridoxine hydrochloride
- Nicotinamide
- Cyanocobalamin
- Iron dextran
- Sodium selenite pentahydrate
- Copper sulphate, anhydrous
- Serum, porcine
- RIBOFLAVIN SODIUM PHOSPHATE

Product identification

Medicine name:
Suibiofer Se (586 mg + 0,998 mg + 0,00403 mg + 0,03 mg + 0,01 mg + 0,0025 mg + 0,08 mg + 0,4 mg)/ml, surowica świńska do 1 ml Roztwór do wstrzykiwań

Active substance:
Thiamine hydrochloride

Pyridoxine hydrochloride

Nicotinamide

Cyanocobalamin

Iron dextran

Sodium selenite pentahydrate

Copper sulphate, anhydrous

Serum, porcine

RIBOFLAVIN SODIUM PHOSPHATE

Target species:

Pig

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Thiamine hydrochloride

0.03 milligram(s) / 1.00 millilitre(s)

Pyridoxine hydrochloride

0.00 milligram(s) / 1.00 millilitre(s)

Nicotinamide

0.40 milligram(s) / 1.00 millilitre(s)

Cyanocobalamin

0.08 milligram(s) / 1.00 millilitre(s)

Iron dextran

586.00 milligram(s) / 1.00 millilitre(s)

Sodium selenite pentahydrate

1.00 milligram(s) / 1.00 millilitre(s)

Copper sulphate, anhydrous

0.00 milligram(s) / 1.00 millilitre(s)

Serum, porcine

1.00 millilitre(s) / 1.00 millilitre(s)

RIBOFLAVIN SODIUM PHOSPHATE
0.01 milligram(s)/millilitre / 1.00 milligram(s)/millilitre

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Pig**
  - Meat and offal. 14 day

Intramuscular use:

- **Pig**
  - Meat and offal. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09A

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

Authorisation status:
Valid

Authorised in:
Poland

Package description:
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Biowet Drwalew Sp. z o.o.

Marketing authorisation date:
19/03/2001

Manufacturing sites for batch release:
Drwalewskie Zaklady Przemyslu Bioweterynaryjnego S.A.

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

Authorisation number:
1121

Date of authorisation status change:
19/03/2001

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.

Summary of Product Characteristics

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.