Selevitam opl. pro inj.

Authorised

- SODIUM SELENITE ANHYDROUS
- DL-ALPHA-TOCOPHEROL

Product identification

Medicine name:
Selevitam opl. pro inj.

Active substance:
SODIUM SELENITE ANHYDROUS
DL-ALPHA-TOCOPHEROL

Target species:
Cattle (calf)
Sheep (lamb)
Pig
Pig (piglet)
Dog

Route of administration:
Intramuscular use

Product details

Active substance and strength:
SODIUM SELENITE ANHYDROUS
0.50 milligram(s) / 1.00 millilitre(s)
DL-ALPHA-TOCOPHEROL

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle (calf)**
  - Meat and offal. 0 day
- **Sheep (lamb)**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Pig (piglet)**
  - Meat and offal. 0 day
- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JC

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
A.A.-Vet Diergeneesmiddelen N.V.

Marketing authorisation date:
19/01/1994

Manufacturing sites for batch release:
A.A.-Vet Diergeneesmiddelen N.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 5123

Date of authorisation status change:
11/02/2013

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

Documents

Combined File of all Documents

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