Selevitan, vet. stungulyf, lausn

Authorised

- Selenide sodium
- DL-ALPHA TOCOPHEROL ACETATE

Product identification

Medicine name:
Selevitan, vet. stungulyf, lausn

Active substance:
Selenide sodium

DL-ALPHA TOCOPHEROL ACETATE

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Selenide sodium
0.60 milligram(s) / 1.00 millilitre(s)

DL-ALPHA TOCOPHEROL ACETATE
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**

  - Meat and offal. 30 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12CE99

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

Authorisation status:
Valid

Authorised in:
Iceland

Package description:
Available only in Icelandic

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Boehringer Ingelheim Animal Health Denmark A/S

Marketing authorisation date:
19/05/1983

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
822959

Date of authorisation status change:
17/02/2011

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.
