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Myogaster-E (100 mg + 1,315 mg)/ml Roztwór do wstrzykiwań

Authorised

- DL-ALPHA TOCOPHEROL ACETATE
- SODIUM SELENITE ANHYDROUS

Product identification

Medicine name:
Myogaster-E (100 mg + 1,315 mg)/ml Roztwór do wstrzykiwań

Active substance:
DL-ALPHA TOCOPHEROL ACETATE
SODIUM SELENITE ANHYDROUS

Target species:
Sheep
Pig
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
DL-ALPHA TOCOPHEROL ACETATE
100.00 milligram(s)/millilitre / 1.00 milligram(s)/millilitre

SODIUM SELENITE ANHYDROUS
1.32 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Sheep**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Cattle**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JB

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

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
V.M.D.

Marketing authorisation date:
24/03/2010

Manufacturing sites for batch release:
VMD N.V.

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

Authorisation number:
1961

Date of authorisation status change:
24/03/2010

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

Documents

Package Leaflet

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Labelling

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.