MYOTROFIL

Authorised

- ADENOSINE PHOSPHATE, SODIUM SALT
- Carnitine hydrochloride
- Sodium selenite
- Magnesium aspartate tetrahydrate
- Potassium aspartate hemihydrate
- Heptaminol hydrochloride

Product identification

Medicine name:
MYOTROFIL

Active substance:
ADENOSINE PHOSPHATE, SODIUM SALT
Carnitine hydrochloride
Sodium selenite
Magnesium aspartate tetrahydrate
Potassium aspartate hemihydrate
Heptaminol hydrochloride

Target species:
Cattle
Dog
Sheep
Cat
Pig

Horse (food producing)

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
ADENOSINE PHOSPHATE, SODIUM SALT
0.25 gram(s) / 100.00 millilitre(s)

Carnitine hydrochloride
0.20 gram(s) / 100.00 millilitre(s)

Sodium selenite
0.05 gram(s) / 100.00 millilitre(s)

Magnesium aspartate tetrahydrate
1.75 gram(s) / 100.00 millilitre(s)

Potassium aspartate hemihydrate
1.30 gram(s) / 100.00 millilitre(s)

Heptaminol hydrochloride
0.50 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Sheep**

- Milk. 0 day
- Meat and offal. 0 day

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

- **Horse (food producing)**
  - Milk. 0 day
  - Meat and offal. 0 day

Intravenous use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day

- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day

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

- **Horse (food producing)**
  - Milk. 0 day
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Milk. 0 hour

- Meat and offal. 0 day

- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Horse (food producing)**
  - Milk. 0 day
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA13A

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian

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:
Izo S.r.l.

Marketing authorisation date:
19/11/1993

Manufacturing sites for batch release:
Izo S.r.l.
Izo S.r.l.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
19/11/1993

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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