CALMOSYL

Authorised

- Strychnos ignatii C7
- Passiflora incarnata C2
- Zincum metallicum C5
- MOSCHUS C5
- Oenanthe crocata C7
- Datura stramonium C5
- HYOSCYAMUS NIGER C5

Product identification

Medicine name:
CALMOSYL

Active substance:
Strychnos ignatii C7

Passiflora incarnata C2

Zincum metallicum C5

MOSCHUS C5

Oenanthe crocata C7

Datura stramonium C5

HYOSCYAMUS NIGER C5

Target species:
Cattle
Pig
Horse

Horse (mare)
Sheep
Goat

Route of administration:
Oral use

Product details

Active substance and strength:
Strychnos ignatii C7
0.14 gram(s) / 1.00 millilitre(s)

Passiflora incarnata C2
0.14 gram(s) / 1.00 millilitre(s)

Zincum metallicum C5
0.14 gram(s) / 1.00 millilitre(s)

MOSCHUS C5
0.14 gram(s) / 1.00 millilitre(s)

Oenanthe crocata C7
0.14 gram(s) / 1.00 millilitre(s)

Datura stramonium C5
0.14 gram(s) / 1.00 millilitre(s)

HYOSCYAMUS NIGER C5
0.14 gram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:
Oral use:

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

- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Horse (mare)**
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV03AX

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French

Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Homeopathic medicinal products (Article 19 of Directive No 2001/82/EC)

Marketing authorisation holder:
Boiron

Marketing authorisation date:
28/06/2012

Manufacturing sites for batch release:
Boiron

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1734122 4/2012

Date of authorisation status change:
21/06/2017

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

Documents

Summary of Product Characteristics

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