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OSTEOTRAUMYL POUDRE ORALE

Authorised

- CALCIUM CARBONICUM HAHNEMANNI D3
- RUTA GRAVEOLENS D3
- SYMPHYTUM OFFICINALE D3
- ARNICA MONTANA D3
- SULFUR IODATUM D3
- CALCAREA PHOSPHORICA D3
- CALCIUM FLUORATUM D3

Product identification

Medicine name:
OSTEOTRAUMYL POUDRE ORALE

Active substance:
CALCIUM CARBONICUM HAHNEMANNI D3
RUTA GRAVEOLENS D3
SYMPHYTUM OFFICINALE D3
ARNICA MONTANA D3
SULFUR IODATUM D3
CALCAREA PHOSPHORICA D3
CALCIUM FLUORATUM D3

Target species:
Cattle
Pig
Equid

Sheep
Goat

Route of administration:
Oral use

Product details

Active substance and strength:
CALCIUM CARBONICUM HAHNEMANNI D3
0.14 gram(s) / 1.00 gram(s)

RUTA GRAVEOLENS D3
0.14 gram(s) / 1.00 gram(s)

SYMPHYTUM OFFICINALE D3
0.14 gram(s) / 1.00 gram(s)

ARNICA MONTANA D3
0.14 gram(s) / 1.00 gram(s)

SULFUR IODATUM D3
0.10 gram(s) / 1.00 gram(s)

CALCAREA PHOSPHORICA D3
0.14 gram(s) / 1.00 gram(s)

CALCIUM FLUORATUM D3
0.10 gram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:
Oral use:

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

- Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

- **Pig**
  - Meat and offal. 0 day
- **Equid**
  - Meat and offal. 0 day
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

- **Sheep**
  - Meat and offal. 0 day
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

- **Goat**
  - Meat and offal. 0 day
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV03AX

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

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Authorisation status:
Valid

Authorised 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:
15/06/2012

Manufacturing sites for batch release:
Boiron

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

Authorisation number:
FR/V/0913144 6/2012

Date of authorisation status change:
31/05/2017

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 and Labelling

This document does not exist in this language (English). You can find it in another language below.