DOLISOVET INTRAMAMMAIRE POMMADE INTRAMAMMAIRE

Authorised

- ATROPA BELLA-DONNA D1
- Solanum dulcamara C1
- ECHINACEA D1
- CALENDULA MOTHER TINCTURE

Product identification

Medicine name:
DOLISOVET INTRAMAMMAIRE POMMADE INTRAMAMMAIRE

Active substance:
ATROPA BELLA-DONNA D1

Solanum dulcamara C1

ECHINACEA D1

CALENDULA MOTHER TINCTURE

Target species:
Cattle (cow)
Sheep (ewe)
Goat (adult female)

Route of administration:
Intramammary use

Product details

Active substance and strength:
ATROPA BELLA-DONNA D1
0.08 gram(s) / 1.00 Syringe

Solanum dulcamara C1
0.05 gram(s) / 1.00 Syringe

ECHINACEA D1
0.08 gram(s) / 1.00 Syringe

CALENDULA MOTHER TINCTURE
0.05 gram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary ointment

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. no withdrawal period

Ne pas livrer le lait du ou des quartiers traités pendant la durée du traitement.

- **Sheep (ewe)**
  - Milk. no withdrawal period

Ne pas livrer le lait du ou des quartiers traités pendant la durée du traitement.

  - Meat and offal. 0 day

- **Goat (adult female)**
  - Milk. no withdrawal period

Ne pas livrer le lait du ou des quartiers traités pendant la durée du traitement.

- Meat and offal. 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
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:
1/03/2006

Manufacturing sites for batch release:
Boiron

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

Authorisation number:
FR/V/7148822 6/2006

Date of authorisation status change:
1/03/2011

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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