Uierbalsem, emulsie voor uitwendig gebruik

Authorised

- LAUREL LEAF OIL
- ROSEMARY OIL
- CLOVE OIL
- ARNICA TINCTURE
- EUCALYPTUS OIL
- Camphor, racemic
- HYPERICI OLEUM

Product identification

Medicine name:
Uierbalsem, emulsie voor uitwendig gebruik

Active substance:
LAUREL LEAF OIL

ROSEMARY OIL

CLOVE OIL

ARNICA TINCTURE

EUCALYPTUS OIL

Camphor, racemic

HYPERICI OLEUM

Target species:
Cattle

Ruminant
Goat (adult female)
Sheep
Equid

Route of administration:
Teat use

Product details

Active substance and strength:
LAUREL LEAF OIL
5.00 milligram(s) / 1.00 gram(s)

ROSEMARY OIL
5.00 milligram(s) / 1.00 gram(s)

CLOVE OIL
0.60 milligram(s) / 1.00 gram(s)

ARNICA TINCTURE
25.00 milligram(s) / 1.00 gram(s)

EUCALYPTUS OIL
20.00 milligram(s) / 1.00 gram(s)

Camphor, racemic
25.00 milligram(s) / 1.00 gram(s)

HYPERICI OLEUM
100.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Cutaneous emulsion

Withdrawal period by route of administration:
Teat use:

- **Cattle**
  - Milk. no withdrawal period nul dagen

- Meat and offal. no withdrawal period nul dagen

- **Ruminant**
  - Milk. no withdrawal period nul dagen
  - Meat and offal. no withdrawal period nul dagen

- **Goat (adult female)**
  - Milk. no withdrawal period nul dagen
  - Meat and offal. no withdrawal period nul dagen

- **Sheep**
  - Milk. no withdrawal period nul dagen
  - Meat and offal. no withdrawal period nul dagen

- **Equid**
  - Meat and offal. no withdrawal period nul dagen

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG52C

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

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Authorisation status:
Valid

Authorised in:
Netherlands

Package description:

Available only in Dutch
Available only in Dutch
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
SaluVet GmbH

Marketing authorisation date:
16/01/1992

Manufacturing sites for batch release:
SaluVet GmbH

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 5627

Date of authorisation status change:
21/10/2020

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

Documents

Combined File of all Documents

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