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JODOFOAM, intrauterinní pěna

Authorised

- Potassium iodate
- Iodine

Product identification

Medicine name:
JODOFOAM, intrauterinní pěna

Active substance:
Potassium iodate

Iodine

Target species:
Cattle (cow)

Route of administration:
Intrauterine use

Product details

Active substance and strength:
Potassium iodate
400.00 milligram(s) / 1.00 Bottle

Iodine
200.00 milligram(s) / 1.00 Bottle

Pharmaceutical form:

Intrauterine foam

Withdrawal period by route of administration:

Intrauterine use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AD30

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

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech

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:
Fortevit Kft.

Marketing authorisation date:
17/07/2000

Manufacturing sites for batch release:
Pernix Pharma Kft.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/051/00-C

Date of authorisation status change:
3/11/2011

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

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

Labelling

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