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JODOFOAM Endofoam 200 mg/400 mg intrauterinná pena

Authorised

- Potassium iodide
- Iodine

Product identification

Medicine name:
JODOFOAM Endofoam 200 mg/400 mg intrauterinná pena

Active substance:
Potassium iodide

Iodine

Target species:
Cattle (cow)

Route of administration:
Intrauterine use

Product details

Active substance and strength:
Potassium iodide
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)**
  - All relevant tissues. 0 day — Without withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AD30

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

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Fortevit Kft.

Marketing authorisation date:

23/12/1997

Manufacturing sites for batch release:
Pernix Pharma Kft.

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

Authorisation number:
96/0639/97-S

Date of authorisation status change:
23/12/1997

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

Documents

Combined File of all Documents

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