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# Jodofoam Endofoam A.U.V. intrauterīnās putas govīm

Authorised

- Iodine
- Potassium iodide

## Product identification

#### **Medicine name:**

Jodofoam Endofoam A.U.V. intrauterīnās putas govīm

#### **Active substance:**

**lodine** 

Potassium iodide

### **Target species:**

Cattle (cow)

#### **Route of administration:**

Intrauterine use

# **Product details**

#### **Active substance and strength:**

**lodine** 

200.00 milligram(s) / 1.00 Vial

Potassium iodide

#### **Pharmaceutical form:**

Intrauterine foam

# Withdrawal period by route of administration: Intrauterine use:

Cattle (cow)

- Meat and offal. 0 day
- Milk. 0 day

#### Anatomical therapeutic chemical veterinary (ATCvet) codes:

**OG51AD30** 

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Latvia

#### Package description:

Available only in Latvian

# Additional information

# **Entitlement type:**

Marketing Authorisation

# Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

# Marketing authorisation holder:

Fortevit Kft.

Marketing authorisation date: 22/10/2008
Manufacturing sites for batch release: Pernix Pharma Kft.
Responsible authority: Food And Veterinary Service
Authorisation number: V/NRP/08/1595
Date of authorisation status change: 22/10/2008
To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>
Documents
Package Leaflet
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Labelling
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Summary of Product Characteristics

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