File downloaded on 2026-11-01
Source URL: https://medicines.health.europa.eu/veterinary/en/600000059880

IBR marker żywy Liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań

Authorised

- Bovine herpesvirus 1, strain GK/D gE gene-deleted, Live

Product identification

Medicine name:
IBR marker żywy Liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań

Active substance:
Bovine herpesvirus 1, strain GK/D gE gene-deleted, Live

Target species:
Cattle

Route of administration:
Nasal use
Intramuscular use

Product details

Active substance and strength:
Bovine herpesvirus 1, strain GK/D gE gene-deleted, Live

Pharmaceutical form:

Lyophilisate for suspension for injection

Withdrawal period by route of administration:

Nasal use:

- **Cattle**
  - Meat and offal. 0 day

Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AD02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Available in:

Poland

Package description:

Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Sp. z o.o.

Marketing authorisation date:
18/09/2012

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2211

Date of authorisation status change:
18/09/2012

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

Documents

Summary of Product Characteristics

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

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.