BioBos IBR marker inact., Suspension for injection

Authorised

- Infectious Bovine rhinotracheitis virus, strain BIO-27, Inactivated

Product identification

Medicine name:
BioBos IBR marker inact., Suspension for injection
BioBos IBR marker inact., injekcinė suspensija galvijams

Active substance:
Infectious Bovine rhinotracheitis virus, strain BIO-27, Inactivated

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Infectious Bovine rhinotracheitis virus, strain BIO-27, Inactivated
1.00 relative potency / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**

  - Milk. 0 hour

  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AA03

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

Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Plastic Vial 1 x 50.0 Dose
Plastic Vial 1 x 25.0 Dose
Glass Vial 1 x 50.0 Dose
Glass Vial 1 x 25.0 Dose
Glass Vial 1 x 5.0 Dose
Glass Vial 10 x 5.0 Dose

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:
Bioveta a.s.

Marketing authorisation date:

11/03/2014

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/14/2217/001-006

Date of authorisation status change:
8/02/2018

Reference member state:
Czechia

Procedure number:
CZ/V/0120/001

Concerned member states:
Bulgaria Estonia Hungary Latvia Lithuania Poland Romania Slovakia Slovenia

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

Documents

RV2217.pdf

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