Bovigal injekčná emulzia pre hovädzí dobytok

Authorised

- Pasteurella multocida, serogroup A, strain PM/7, Inactivated
- Bovine parainfluenza 3, strain Mensik, Inactivated
- Bovine herpesvirus 1, strain N3760, Inactivated

Product identification

Medicine name:
Bovigal injekčná emulzia pre hovädzí dobytok

Active substance:
Pasteurella multocida, serogroup A, strain PM/7, Inactivated

Bovine parainfluenza 3, strain Mensik, Inactivated

Bovine herpesvirus 1, strain N3760, Inactivated

Target species:
Cattle
Cattle (calf)

Route of administration:
Intramuscular use
Intramuscular use

Product details

Active substance and strength:

Pasteurella multocida, serogroup A, strain PM/7, Inactivated
1.00 log2 haemagglutination inhibiting unit(s) / 3.00 millilitre(s)

Bovine parainfluenza 3, strain Mensik, Inactivated
5.00 log2 haemagglutination inhibiting unit(s) / 3.00 millilitre(s)

Bovine herpesvirus 1, strain N3760, Inactivated
1.00 log2 virus neutralising unit(s) / 3.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - All relevant tissues. 0 day

Intramuscular use:

- **Cattle (calf)**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AV

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak

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:
Pharmagal Bio spol. s r.o.

Marketing authorisation date:
9/06/2003

Manufacturing sites for batch release:
Pharmagal Bio spol. s r.o.

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

Authorisation number:
97/028/03-S

Date of authorisation status change:
29/11/2022

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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