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BioBos Respi 4 injekčná suspenzia pre hovädzí dobytok

Authorised

- Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated
- Bovine viral diarrhoea virus, strain BIO-25, Inactivated
- Bovine parainfluenza virus 3, strain BIO-23, Inactivated
- Bovine respiratory syncytial virus, strain BIO-24, Inactivated

Product identification

Medicine name:
BioBos Respi 4 injekčná suspenzia pre hovädzí dobytok

Active substance:
Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated

Bovine viral diarrhoea virus, strain BIO-25, Inactivated

Bovine parainfluenza virus 3, strain BIO-23, Inactivated

Bovine respiratory syncytial virus, strain BIO-24, Inactivated

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Bovine viral diarrhoea virus, strain BIO-25, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Bovine parainfluenza virus 3, strain BIO-23, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Bovine respiratory syncytial virus, strain BIO-24, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Available only in Slovak

Available only in Slovak

Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
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Available only in Slovak

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:
Vetservis s.r.o.

Marketing authorisation date:
27/06/2013

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
97/043/13-S

Date of authorisation status change:
27/06/2013

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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