BioBos Respi 4 vakcina A.U.V.

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 vakcina A.U.V.

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 / 1.00 unit(s)/dose

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

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

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

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Hungary

Package description:

Available only in Hungarian

Available only in Hungarian

Available only in Hungarian
Available only in Hungarian
Available only in Hungarian
Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Alphavet Zrt.

Marketing authorisation date:
7/12/2016

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
7/12/2016

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