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Bovipast RSP vakcina A.U.V.

Authorised

- Bovine parainfluenza virus 3, strain SF-4, Inactivated
- Bovine respiratory syncytial virus, strain EV 908, Inactivated
- Mannheimia haemolytica, Inactivated

Product identification

Medicine name:
Bovipast RSP vakcina A.U.V.

Active substance:
Bovine parainfluenza virus 3, strain SF-4, Inactivated

Bovine respiratory syncytial virus, strain EV 908, Inactivated

Mannheimia haemolytica, Inactivated

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Bovine parainfluenza virus 3, strain SF-4, Inactivated
7.30 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

Bovine respiratory syncytial virus, strain EV 908, Inactivated
5.50 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

Mannheimia haemolytica, Inactivated
9000.00 million cells / 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:
QI02AL04

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

Authorisation status:
Valid

Authorised in:
Hungary

Available in:
Hungary

Package description:
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:
Intervet International B.V.

Marketing authorisation date:
9/10/2003

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Directorate Of Veterinary Medicinal Products

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

Date of authorisation status change:
9/10/2003

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