Pneumovac Suspension for Injection for Cattle

Authorised

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

Product identification

Medicine name:
Pneumovac suspension for injection for cattle
Pneumovac Suspension for Injection for Cattle

Active substance:
Bovine respiratory syncytial virus, strain BIO-24, Inactivated

Bovine parainfluenza virus 3, strain BIO-23, Inactivated

Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Bovine respiratory syncytial virus, strain BIO-24, Inactivated
1.00 relative potency / 1.00 Dose

Bovine parainfluenza virus 3, strain BIO-23, Inactivated
1.00 relative potency / 1.00 Dose

Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated
1.00 relative potency / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

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

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:
United Kingdom (Northern Ireland)

Package description:
The vaccine is filled in glass vials:Hydrolytic type I 10ml vials containing 10 ml (5 doses). All containers are closed with chlorobutyl rubber stoppers and secured with aluminium seals. The product is delivered as follows: 10 x 10ml in a transparent plastic box with cover. Carton for mass packaging 10 x 10 mlEach package contains an approved Package Leaflet
In plastic vials:15ml vials containing 10 ml (5 doses)All containers are closed with chlorobutyl rubber stoppers and secured with aluminium seals.The product is

delivered as follows:1 x 10ml in cardboard boxesEach package contains an approved Package Leaflet
The vaccine is filled in glass vials:100ml vials containing 100 ml (50 doses). All containers are closed with chlorobutyl rubber stoppers and secured with aluminium seals. The product is delivered as follows:1 x 100ml in cardboard boxes. Each package contains an approved Package Leaflet.
The vaccine is filled in glass vials:Hydrolytic type I 50ml vials containing 50 ml (25 doses). All containers are closed with chlorobutyl rubber stoppers and secured with aluminium seals. The product is delivered as follows:1 x 50ml in cardboard boxes. Each package contains an approved Package Leaflet.
The vaccine is filled in glass vials:Hydrolytic type I 10ml vials containing 10 ml (5 doses). All containers are closed with chlorobutyl rubber stoppers and secured with aluminium seals. The product is delivered as follows:1 x 10ml in cardboard boxes. Each package contains an approved Package Leaflet.
In plastic vials:60ml vials containing 50ml (25 doses)All containers are closed with chlorobutyl rubber stoppers and secured with aluminium seals.The product is delivered as follows:1 x 50ml in cardboard boxesEach package contains an approved Package Leaflet
In plastic vials:120ml vials containing 100ml (50 doses)All containers are closed with chlorobutyl rubber stoppers and secured with aluminium seals.The product is delivered as follows:1 x 100ml in cardboard boxesEach package contains an approved Package Leaflet
In plastic vials:15ml vials containing 10 ml (5 doses)All containers are closed with chlorobutyl rubber stoppers and secured with aluminium seals.The product is delivered as follows: 10 x 10ml in a transparent plastic box with cover. Carton for mass packaging 10 x 10 mlEach package contains an approved Package Leaflet

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:
Animal Health Distributors Limited

Marketing authorisation date:
17/12/2020

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 51609/4000

Date of authorisation status change:
24/08/2022

Reference member state:
Ireland

Procedure number:
IE/V/0638/001

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics
