Bovilis IBR marker inac suspension for injection for cattle

Not authorised

- Bovine herpesvirus 1, strain GK/D gE gene-deleted, Inactivated

Product identification

Medicine name:
Bovilis IBR marker inac suspension for injection for cattle

Active substance:
Bovine herpesvirus 1, strain GK/D gE gene-deleted, Inactivated

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Bovine herpesvirus 1, strain GK/D gE gene-deleted, Inactivated
60.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AA03

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

Authorisation status:
Surrendered

Authorised in:
Greece

Package description:
(ID27) 2000 millilitre(s): Box (Cardboard) with 10 Bottle (PolyEthylene TerePhthalate) each with 200 millilitre(s)
(ID26) 2000 millilitre(s): Box (Cardboard) with 10 Bottle (Glass) each with 200 millilitre(s)
(ID25) 200 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 200 millilitre(s)
(ID24) 200 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 200 millilitre(s)
(ID23) 1000 millilitre(s): Box (Cardboard) with 10 Bottle (PolyEthylene TerePhthalate) each with 100 millilitre(s)
(ID22) 1000 millilitre(s): Box (Cardboard) with 10 Bottle (Glass) each with 100 millilitre(s)
(ID21) 100 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 100 millilitre(s)
(ID20) 100 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 100 millilitre(s)

(ID19) 500 millilitre(s): Box (Cardboard) with 10 Bottle (PolyEthylene TerePhthalate) each with 50 millilitre(s)
(ID17) 500 millilitre(s): Box (Cardboard) with 10 Bottle (Glass) each with 50 millilitre(s)
(ID15) 50 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 50 millilitre(s)
(ID10) 50 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 50 millilitre(s)
(ID9) 200 millilitre(s): Box (Cardboard) with 10 Bottle (PolyEthylene TerePhthalate) each with 20 millilitre(s)
(ID8) 200 millilitre(s): Box (Cardboard) with 10 Bottle (Glass) each with 20 millilitre(s)
(ID7) 20 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 20 millilitre(s)
(ID6) 20 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 20 millilitre(s)
(ID4) 100 millilitre(s): Box (Cardboard) with 10 Bottle (PolyEthylene TerePhthalate) each with 10 millilitre(s)
(ID3) 100 millilitre(s): Box (Cardboard) with 10 Bottle (Glass) each with 10 millilitre(s)
(ID2) 10 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 10 millilitre(s)
(ID1) 10 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 10 millilitre(s)

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:
3/12/2006

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

Responsible authority:
National Organization For Medicines

Authorisation number:
81989/23-11-2011/K-0163501

Date of authorisation status change:
20/05/2024

Reference member state:
Germany

Procedure number:
DE/V/0237/001

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