Rispoval IBR-Marker Vivum, Lyophilisate and diluent for suspension for injection for cattle

Authorised

- Bovine herpesvirus 1, strain Difivac gE gene-deleted, Live

Product identification

Medicine name:
Rispoval IBR-Marker Live
Rispoval IBR-Marker Vivum, Lyophilisate and diluent for suspension for injection for cattle

Active substance:
Bovine herpesvirus 1, strain Difivac gE gene-deleted, Live

Target species:
Cattle
Cattle (for meat production)
Cattle (calf)
Cattle (heifer)
Cattle (suckling calf)

Route of administration:
Nasal use
Nasal use

Product details

Active substance and strength:
Bovine herpesvirus 1, strain Difivac gE gene-deleted, Live
10000000.00 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:
Nasal use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Cattle (for meat production)**
  - Meat and offal. 0 day
- **Cattle (calf)**
  - Meat and offal. 0 day
- **Cattle (heifer)**
  - Meat and offal. 0 day
- **Cattle (suckling calf)**
  - Meat and offal. 0 day

Nasal use:

- **Cattle**
  - Milk. 0 day

- Meat and offal. 0 day

- **Cattle (for meat production)**
  - Meat and offal. 0 day

- **Cattle (calf)**
  - Meat and offal. 0 day

- **Cattle (heifer)**
  - Meat and offal. 0 day

- **Cattle (suckling calf)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AD01

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
(ID2) 50 Dose; 100 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 50 Dose and 1 Bottle (Glass) with 100 millilitre(s)
(ID1) 10 Dose; 20 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 10 Dose and 1 Bottle (Glass) with 20 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:
Zoetis Belgium S.A.

Marketing authorisation date:
9/12/2013

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10387/062/001

Date of authorisation status change:
9/12/2013

Reference member state:
Germany

Procedure number:
DE/V/0022/001

Concerned member states:
Belgium Bulgaria Czechia Estonia Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics