Bovilis Ringvac <9 and >21 Lyophilisate+solvent for suspension for injection

Authorised

- Trichophyton verrucosum, strain LTF-130, Live

Product identification

Medicine name:
Bovilis Ringvac <9 and >21 Lyophilisate+solvent for suspension for injection
Bovilis Ringvac lyofilizát a rozpúšťadlo pre injekčnú suspenziu pre hovädzí dobytok

Active substance:
Trichophyton verrucosum, strain LTF-130, Live

Target species:
Cattle
Cattle (calf)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Trichophyton verrucosum, strain LTF-130, Live
21000000.00 cells / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AP01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

(ID2) 40 millilitre(s); 40 Dose: Box (Cardboard) with 1 Bottle (Glass) with 40 Dose and 1 Bottle (Glass) with 40 millilitre(s)
(ID1) 10 Dose; 10 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 10 Dose and 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:
This information is not available for this product.

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/004/MR/05-S

Date of authorisation status change:
1/03/2005

Reference member state:
Germany

Procedure number:
DE/V/0231/001

Concerned member states:
Belgium Bulgaria Cyprus Czechia France Greece Hungary Ireland Luxembourg Poland Portugal Slovakia Slovenia

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

Documents

Summary of Product Characteristics

Combined File of all Documents

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