Bovilis Ringvac lyophilisate and solvent for suspension for injection for cattle

Not authorised

- Trichophyton verrucosum, strain LTF-130, Live

Product identification

Medicine name:
Bovilis Ringvac lyophilisate and solvent for suspension for injection for cattle

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:
Surrendered

Authorised in:
Ireland

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 (Ireland) Limited

Marketing authorisation date:
21/04/2017

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/284/001

Date of authorisation status change:
7/10/2025

Reference member state:
Germany

Procedure number:
DE/V/0231/001

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