

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 vakcina A.U.V.

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:**

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Cattle

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

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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:

Hungary

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:

15/05/2017

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

3877/X/17 NÉBIH ÁTI

Date of authorisation status change:

15/05/2017

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