BOVILIS RINGVAC, liofilizat i otapalo za suspenziju za injekciju, za goveda

Not authorised

- Trichophyton verrucosum, strain LTF-130, Live

Product identification

Medicine name:
BOVILIS RINGVAC, liofilizat i otapalo za suspenziju za injekciju, za goveda

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

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Trichophyton verrucosum, strain LTF-130, Live
21000000.00 Piece / 1.00 millilitre(s)

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AP01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Croatia

Package description:

Available only in Croatian

Available only in Croatian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V. Subsidiary In The Republic Of Croatia

Marketing authorisation date:

21/09/2015

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

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/13-01/172

Date of authorisation status change:
23/02/2022

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

Documents

Summary of Product Characteristics

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