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Fixr trichoben, Lyophilisate and solvent for suspension for injection for cattle

Authorised

- Trichophyton verrucosum, strain Bodin 1902, Live

Product identification

Medicine name:
Fixr trichoben, Lyophilisate and solvent for suspension for injection for cattle

Active substance:
Trichophyton verrucosum, strain Bodin 1902, Live

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Trichophyton verrucosum, strain Bodin 1902, Live
6.00 unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 14 day
  - Meat and offal. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02A

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Package description:

Carton box containing 1 x 80 ml (after reconstitution) and corresponding amounts of Diluent A

Carton box containing 5 x 10 ml (after reconstitution) and corresponding amounts of Diluent A

Carton box containing 1 x 40 ml (after reconstitution) and corresponding amounts of Diluent A

Diluent A is supplied in vials of hydrolytic quality II sealed with rubber stopper and aluminium cap resentation 40 ml in 50-ml vial

Vaccine is supplied in the freeze-dried state in 10-ml vials, hydrolytic quality I, sealed with rubber stopper and aluminium Flip off

Vaccine is supplied in the freeze-dried state in 10-ml vials, hydrolytic quality I, sealed with rubber stopper and aluminium cap

Diluent A is supplied in vials of hydrolytic quality I sealed with rubber stopper and aluminium cap presentation 10 ml in 10-ml vial

Diluent A is supplied in vials of hydrolytic quality I sealed with rubber stopper and aluminium cap presentation 80 ml in 100-ml vial
Diluent A is supplied in vials of hydrolytic quality I sealed with rubber stopper and aluminium cap resentation 40 ml in 50-ml vial
Diluent A is supplied in vials of hydrolytic quality II sealed with rubber stopper and aluminium cap presentation 10 ml in 10-ml vial
Diluent A is supplied in vials of hydrolytic quality II sealed with rubber stopper and aluminium cap presentation 80 ml in 100-ml vial

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Kernfarm B.V.

Marketing authorisation date:
9/04/2018

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
9/04/2018

Reference member state:
Netherlands

Procedure number:

NL/V/0233/001

Concerned member states:
Belgium

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

Documents

Combined File of all Documents

Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.