SILIRUM EMULSION INJECTABLE POUR BOVINS

Authorised

- Mycobacterium avium, subsp. paratuberculosis, strain 316F, Inactivated

Product identification

Medicine name:
SILIRUM EMULSION INJECTABLE POUR BOVINS

Active substance:
Mycobacterium avium, subsp. paratuberculosis, strain 316F, Inactivated

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Mycobacterium avium, subsp. paratuberculosis, strain 316F, Inactivated
1.00 relative unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
CZ Vaccines S.A.U.

Marketing authorisation date:
4/02/2014

Manufacturing sites for batch release:
CZ Vaccines S.A.U.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/0984602 5/2014

Date of authorisation status change:
11/01/2019

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

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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