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B-19 CZV SUBCUTANEA LIOFILIZADO Y DISOLVENTE PARA SUSPENSION INYECTABLE PARA BOVINO

Authorised

- Brucella abortus, strain B-19, Live

Product identification

Medicine name:
B-19 CZV SUBCUTANEA LIOFILIZADO Y DISOLVENTE PARA SUSPENSION INYECTABLE PARA BOVINO

Active substance:
Brucella abortus, strain B-19, Live

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Brucella abortus, strain B-19, Live

10.00 colony forming unit(s)/gram / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AE

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:

CZ Vaccines S.A.U.

Marketing authorisation date:
11/01/1979

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2940 ESP

Date of authorisation status change:
17/12/2013

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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