CUNIVAK PAST

Not authorised

- Bordetella bronchiseptica, Inactivated

Product identification

Medicine name:
CUNIVAK PAST

Active substance:
Bordetella bronchiseptica, Inactivated

Target species:
Rabbit

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Bordetella bronchiseptica, Inactivated
12000000000.00 Organisms / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI08AB01

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

Authorisation status:
Surrendered

Authorised in:
Germany

Package description:
Available only in German
Available only in German
Available only in German
Available only in German

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:
Ceva Tiergesundheit GmbH

Marketing authorisation date:
1/04/1997

Manufacturing sites for batch release:
IDT Biologika GmbH

Responsible authority:
PEI

Authorisation number:
4a/96

Date of authorisation status change:
16/04/2021

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.