BOVISALORAL

Not authorised

- Salmonella enterica, subsp. enterica, serovar Dublin, Live

Product identification

Medicine name:
BOVISALORAL

Active substance:
Salmonella enterica, subsp. enterica, serovar Dublin, Live

Target species:
Cattle

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Dublin, Live
500000000000.00 Colony forming unit / 1.00 Dose

Pharmaceutical form:
Oral lyophilisate

Withdrawal period by route of administration:

In drinking water/milk use:

- **Cattle**
  - Meat and offal. 3 week
  - Milk. 3 week

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AE02

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

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:
3/01/1995

Manufacturing sites for batch release:
IDT Biologika GmbH

Responsible authority:
PEI

Authorisation number:
573a/91

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.