Bovilis BTV8 (WD) (--) - Suspension for injection

Not authorised

- Bluetongue virus, serotype 8, Inactivated

Product identification

Medicine name:
Bovilis BTV8 (WD) (--) - Suspension for injection

Active substance:
Bluetongue virus, serotype 8, Inactivated

Target species:
Cattle
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Bluetongue virus, serotype 8, Inactivated

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
- **Sheep**
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days
  - Not applicable. 0 day Zero days

- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days
- Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AA02

Legal status of supply:
Medicinal product subject to medical prescription

Authorisation status:
Surrendered

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Additional information

Entitlement type:

Marketing Authorisation

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

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
6/09/2010

Manufacturing sites for batch release:
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
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Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.
Intervet International B.V.

Responsible authority:
European Commission

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

Date of authorisation status change:
6/09/2010

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

Documents

Combined File of all Documents

English (PDF)
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