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Scourguard 3

Authorised

- Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), strain NADC 1471, Inactivated
- Bovine rotavirus A, strain Lincoln, Live
- Bovine coronavirus, strain Hansen, Live

Product identification

Medicine name:
Scourguard 3

Active substance:
Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), strain NADC 1471, Inactivated

Bovine rotavirus A, strain Lincoln, Live

Bovine coronavirus, strain Hansen, Live

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), strain NADC 1471, Inactivated
4.50 log2 antibody titre / 1.00 Dose

Bovine rotavirus A, strain Lincoln, Live
10000000.00 50% tissue culture infectious dose / 1.00 Dose

Bovine coronavirus, strain Hansen, Live
100000.00 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AI01

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

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Authorisation status:
Valid

Authorised in:
Germany

Available 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:
Zoetis Deutschland GmbH

Marketing authorisation date:
22/11/2004

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
89a/82

Date of authorisation status change:
29/10/2009

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

Documents

Combined File of all Documents

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