Locatim inj.

Authorised

- Immunoglobulins against Bovine coronavirus, Bovine
- Immunoglobulins against Bovine rotavirus, Bovine
- Immunoglobulins against Escherichia coli F5 antigen, Bovine
- Immunoglobulins against Escherichia coli 078:80B, Bovine
- BOVINE ROTAVIRUS IMMUNOGLOBULIN

Product identification

Medicine name:
Locatim inj.

Active substance:
Immunoglobulins against Bovine coronavirus, Bovine

Immunoglobulins against Bovine rotavirus, Bovine

Immunoglobulins against Escherichia coli F5 antigen, Bovine

Immunoglobulins against Escherichia coli 078:80B, Bovine

BOVINE ROTAVIRUS IMMUNOGLOBULIN

Target species:
Cattle (pre-ruminant)

Route of administration:
Oral use

Subcutaneous use

Product details

Active substance and strength:

Immunoglobulins against Bovine coronavirus, Bovine
0.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Immunoglobulins against Bovine rotavirus, Bovine
0.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Immunoglobulins against Escherichia coli F5 antigen, Bovine
0.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Immunoglobulins against Escherichia coli 078:80B, Bovine
0.01 slow agglutination test unit(s) / 1.00 millilitre(s)

BOVINE ROTAVIRUS IMMUNOGLOBULIN
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Oral use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AM05

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Germany

Package description:
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Melchior Sante Animale

Marketing authorisation date:
14/06/2008

Manufacturing sites for batch release:
Biokema Europe

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.03587.01.1

Date of authorisation status change:
30/01/2013

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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