Decivac FMD DOE

Authorised

- Foot-and-mouth disease virus, serotype A, Inactivated
- Foot-and-mouth disease virus, serotype O, Inactivated
- Foot-and-mouth disease virus, serotype C, Inactivated
- Foot-and-mouth disease virus, serotype Asia 1, Inactivated
- Foot-and-mouth disease virus, serotype SAT 1, Inactivated
- Foot-and-mouth disease virus, serotype SAT 2, Inactivated

Product identification

Medicine name:
Decivac FMD DOE

Active substance:
Foot-and-mouth disease virus, serotype A, Inactivated
Foot-and-mouth disease virus, serotype O, Inactivated
Foot-and-mouth disease virus, serotype C, Inactivated
Foot-and-mouth disease virus, serotype Asia 1, Inactivated
Foot-and-mouth disease virus, serotype SAT 1, Inactivated
Foot-and-mouth disease virus, serotype SAT 2, Inactivated

Target species:
Sheep
Cattle
Goat
Pig

Route of administration:

Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Foot-and-mouth disease virus, serotype A, Inactivated
3.00 50% Protective Dose / 1.00 Dose

Foot-and-mouth disease virus, serotype O, Inactivated
3.00 50% Protective Dose / 1.00 Dose

Foot-and-mouth disease virus, serotype C, Inactivated
3.00 50% Protective Dose / 1.00 Dose

Foot-and-mouth disease virus, serotype Asia 1, Inactivated
3.00 50% Protective Dose / 1.00 Dose

Foot-and-mouth disease virus, serotype SAT 1, Inactivated
3.00 50% Protective Dose / 1.00 Dose

Foot-and-mouth disease virus, serotype SAT 2, Inactivated
3.00 50% Protective Dose / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

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

Goat

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

•

Pig

- Meat and offal. 0 day

Subcutaneous use:

•

Sheep

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

•

Pig

- Meat and offal. 0 day

•

Goat

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

•

Cattle

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AA04

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
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:
Intervet Deutschland GmbH

Marketing authorisation date:
3/08/2000

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
BFAV/MKS/2/2000

Date of authorisation status change:
9/08/2010

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.