CLASOVAX vaccino inattivato in sospensione iniettabile per bovini, ovini e caprini

Authorised

- Salmonella enterica, subsp. enterica, serovar Abortusovis, strain ATCC 31684, Inactivated
- Chlamydia abortus, strain AB7, Inactivated
- Chlamydia abortus, Inactivated
- Aluminium hydroxide

Product identification

Medicine name:
CLASOVAX vaccino inattivato in sospensione iniettabile per bovini, ovini e caprini

Active substance:
Salmonella enterica, subsp. enterica, serovar Abortusovis, strain ATCC 31684, Inactivated

Chlamydia abortus, strain AB7, Inactivated

Chlamydia abortus, Inactivated

Aluminium hydroxide

Target species:
Cattle
Goat
Sheep

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Abortusovis, strain ATCC 31684, Inactivated
640.00 slow agglutination test unit(s) / 2.00 millilitre(s)

Chlamydia abortus, strain AB7, Inactivated
200.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Chlamydia abortus, Inactivated
200.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Aluminium hydroxide
0.25 millilitre(s) / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

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

Subcutaneous use:

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

- **Sheep**

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AB

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian

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:
MSD Animal Health S.r.l.

Marketing authorisation date:
4/09/2000

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:

Ministry Of Health

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

Date of authorisation status change:
4/09/2000

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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