SYVA BAX

Authorised

- Clostridium perfringens, type A, alpha toxoid
- Clostridium perfringens, type B and C, beta toxoid
- Clostridium perfringens, type B and D, epsilon toxoid
- Clostridium septicum, toxoid
- Clostridium novyi, type B, alpha toxoid
- Clostridium tetani, toxoid
- Clostridium sordellii, toxoid
- Clostridium chauvoei, Inactivated

Product identification

Medicine name:
СИВА БАКС
SYVA BAX

Active substance:
Clostridium perfringens, type A, alpha toxoid
Clostridium perfringens, type B and C, beta toxoid
Clostridium perfringens, type B and D, epsilon toxoid
Clostridium septicum, toxoid
Clostridium novyi, type B, alpha toxoid
Clostridium tetani, toxoid
Clostridium sordellii, toxoid
Clostridium chauvoei, Inactivated

Target species:
Cattle
Pig

Sheep
Goat
Rabbit

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Clostridium perfringens, type A, alpha toxoid
0.30 international unit(s) / 2.00 millilitre(s)

Clostridium perfringens, type B and C, beta toxoid
10.00 international unit(s) / 2.00 millilitre(s)

Clostridium perfringens, type B and D, epsilon toxoid
10.00 international unit(s) / 2.00 millilitre(s)

Clostridium septicum, toxoid
2.50 international unit(s) / 2.00 millilitre(s)

Clostridium novyi, type B, alpha toxoid
3.50 international unit(s) / 2.00 millilitre(s)

Clostridium tetani, toxoid
2.50 international unit(s) / 2.00 millilitre(s)

Clostridium sordellii, toxoid
100.00 100% protective dose / 2.00 millilitre(s)

Clostridium chauvoei, Inactivated
100.00 100% protective dose / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:
- **Cattle**
- **Pig**

Subcutaneous use:

- **Cattle**
- **Sheep**
- **Goat**
- **Rabbit**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB
QI03AB
QI04AB
QI08AB
QI09AB

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Laboratorios Syva S.A.

Marketing authorisation date:
3/02/2008

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

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1858

Date of authorisation status change:
20/09/2012

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

Documents

Summary of Product Characteristics

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Package Leaflet

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