

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, cells and toxin, Inactivated

Product identification

Medicine name:

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, cells and toxin, 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, cells and toxin, Inactivated
100.00 100% protective dose / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

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.

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

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.