

TOXIPRA – S7

Authorised

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

Product identification

Medicine name:

ТОКСИПРА-С7

TOXIPRA – S7

Active substance:

Clostridium sordellii, toxoid

Clostridium chauvoei, Inactivated

Clostridium septicum, toxoid

Clostridium perfringens, type B, C and D, beta toxoid

Clostridium novyi, type B, alpha toxoid

Target species:

Cattle

Sheep

Goat

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Clostridium sordellii, toxoid

0.18 millilitre(s) / 1.00 millilitre(s)

Clostridium chauvoei, Inactivated

0.10 millilitre(s) / 1.00 millilitre(s)

Clostridium septicum, toxoid

3.50 international unit(s) / 1.00 millilitre(s)

Clostridium perfringens, type B, C and D, beta toxoid

10.00 international unit(s) / 1.00 millilitre(s)

Clostridium novyi, type B, alpha toxoid

3.50 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cattle

-

Sheep

-

Goat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised 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 Hipra S.A.

Marketing authorisation date:

24/06/2008

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-1968-06.03.2013

Date of authorisation status change:

5/03/2013

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 and Labelling

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