Suiseng Suspension for injection for pigs

Not authorised

- Escherichia coli, fimbrial adhesin F4ac
- Escherichia coli, fimbrial adhesin F5
- Clostridium perfringens, type C, beta toxoid
- Clostridium novyi, type B, alpha toxoid
- Escherichia coli, fimbrial adhesin F4ab
- Escherichia coli, fimbrial adhesin F6
- Escherichia coli, LT toxoid

Product identification

Medicine name:
Suiseng Suspension for injection for pigs
SUISENG инжекционна суспензия за свине

Active substance:
Escherichia coli, fimbrial adhesin F4ac
Escherichia coli, fimbrial adhesin F5
Clostridium perfringens, type C, beta toxoid
Clostridium novyi, type B, alpha toxoid
Escherichia coli, fimbrial adhesin F4ab
Escherichia coli, fimbrial adhesin F6
Escherichia coli, LT toxoid

Target species:

Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Escherichia coli, fimbrial adhesin F4ac
78.00 percent / 1.00 Dose

Escherichia coli, fimbrial adhesin F5
79.00 percent / 1.00 Dose

Clostridium perfringens, type C, beta toxoid
35.00 percent / 1.00 Dose

Clostridium novyi, type B, alpha toxoid
50.00 percent / 1.00 Dose

Escherichia coli, fimbrial adhesin F4ab
65.00 percent / 1.00 Dose

Escherichia coli, fimbrial adhesin F6
80.00 percent / 1.00 Dose

Escherichia coli, LT toxoid
55.00 percent / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB08

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

Authorisation status:
Surrendered

Authorised in:
Bulgaria

Package description:
Cardboard box with 1 PET vial of 125 doses (250 ml). 250 ml PET plastic vials, closed with type I rubber stoppers and aluminium caps.
Cardboard box with1 PET vial of 50 doses (100 ml). 100 ml PET plastic vials, closed with type I rubber stoppers and aluminium caps.
Cardboard box with 1 glass vial of 25 doses (50 ml). 50 ml Type-I colourless glass vials, closed with type I rubber stoppers and aluminium caps.
Cardboard box with 1 glass vial of 10 doses (20 ml).20 ml Type-I colourless glass vials, closed with type I rubber stoppers and aluminium caps.
Cardboard box with1 Glass vial of 50 doses (100 ml). 100 ml Type-I colourless glass vials, closed with type I rubber stoppers and aluminium caps.
Cardboard box with 1 PET vial of 25 doses (50 ml). , 50 ml PET plastic vials, closed with type I rubber stoppers and aluminium caps
Cardboard box with 1 PET vial of 10 doses (20 ml).20 ml PET plastic vials, closed with type I rubber stoppers and aluminium caps.

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:
Laboratorios Hipra S.A.

Marketing authorisation date:
17/08/2009

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

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2381

Date of authorisation status change:
30/03/2025

Reference member state:
Spain

Procedure number:
ES/V/0461/001

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