BIOSUIS ParvoEry, Suspension for injection

Authorised

- Porcine parvovirus, strain CAPM V198 S-27, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

Product identification

Medicine name:
BIOSUIS ParvoEry, Suspension for injection
BIOSUIS ParvoEry, suspenzija za injekciju, za svinje

Active substance:
Porcine parvovirus, strain CAPM V198 S-27, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

Target species:
Pig (sow)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine parvovirus, strain CAPM V198 S-27, Inactivated
4.00 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

1.00 relative potency / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig (sow)**

  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AL01

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

Authorisation status:
Valid

Authorised in:
Croatia

Package description:
Glass Vial 1 x 10.0 millilitre(s)
Glass Vial 10 x 10.0 millilitre(s)
Glass Vial 1 x 50.0 millilitre(s)
Plastic Vial 1 x 50.0 millilitre(s)
Glass Vial 1 x 100.0 millilitre(s)
Plastic Vial 1 x 100.0 millilitre(s)
Plastic Vial 1 x 250.0 millilitre(s)

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:
Bioveta a.s.

Marketing authorisation date:
1/04/2019

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/19-01/149

Date of authorisation status change:
22/09/2021

Reference member state:
Czechia

Procedure number:
CZ/V/0148/001

Concerned member states:
Belgium Bulgaria Croatia Denmark Estonia Germany Greece Hungary Ireland Latvia Lithuania Netherlands Norway Poland Portugal Spain Sweden United Kingdom (Northern Ireland)

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.
